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(71) Applicant (for all designated States except US): FOAMIX LTD. [IL/IL]; 2 Hashaked Street, 74104 Ness Ziona (IL).

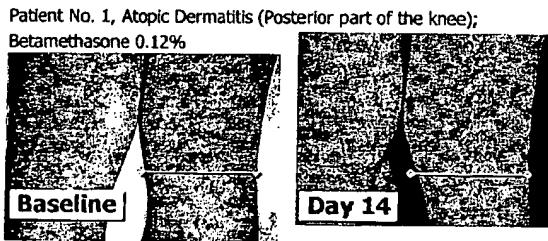
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(72) Inventors; and

(75) Inventors/Applicants (for US only): TAMARKIN, Dov [IL/IL]; 537 Har Hila Street, Maccabim (IL). FRIEDMAN, Doron [IL/IL]; 33 Alon Street, Karmei-Yosef (IL). EINI, Meir [IL/IL]; 2 Hashaked Street, Ness Ziona (IL).

(54) Title: COSMETIC AND PHARMACEUTICAL FOAM

Picture No. 2: Improvement in the treatment of atopic dermatitis, using
Betamethasone 0.12% foam



Patient No. 2, Atopic Dermatitis (Arms); Betamethasone 0.12%



(57) Abstract: The invention relates to an alcohol-free cosmetic or pharmaceutical foam carrier comprising water, a hydrophobic solvent, a foam adjuvant agent, a surface-active agent and a water gelling agent. The cosmetic or pharmaceutical foam carrier does not contain aliphatic alcohols, making it non-irritating and non-drying. The alcohol-free foam carrier is suitable for inclusion of both water-soluble and oil soluble pharmaceutical and cosmetic agents.

WO 2004/037225 A2

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COSMETIC AND PHARMACEUTICAL FOAM

This application claims priority to Related Applications of Israeli patent application No. 152486 filed October 25, 2002 entitled "Alcohol-free Cosmetic and Pharmaceutical Foam Carrier", and to U.S. provisional patent application No. 60/495,546, filed November 29, 2002, entitled "Cosmetic and Pharmaceutical Foam", which are incorporated herein in their entirety.

FIELD OF THE INVENTION

10 The invention relates to an alcohol-free, cosmetic or pharmaceutical foam carrier and its use. More specifically, the invention relates to a cosmetic or pharmaceutical foam carrier suitable for inclusion of both water soluble and oil soluble pharmaceutical and cosmetic agents.

15 BACKGROUND OF THE INVENTION

External topical administration is an important route for the administration of drugs in disease treatment. In external topical administration, the drug is absorbed into and/or through skin, mucous membrane or wound tissue. Many groups of drugs, including, for example, antibiotic, anti-fungal, anti-inflammatory, 20 anesthetic, analgesic, anti-allergic, corticosteroid, retinoid and anti-proliferative medications are preferably administered in hydrophobic media, e.g. ointments or oils. However, due to the undesirable consistency of these hydrophobic carriers, their use is limited. For instance, ointments containing white petrolatum, e.g., Vaseline petroleum jelly, as the carrier often form an impermeable barrier, so that 25 metabolic products and excreta from the wounds to which they are applied are not easily removed or drained away. Furthermore, it is difficult for the active drug dissolved in the carrier to pass through the white petrolatum barrier layer into the wound tissue, so the efficacy of the drug is reduced.

In addition, ointments and creams often do not create an environment for 30 promoting respiration of the wound tissue and it is not favorable to the normal respiration of the skin. An additional disadvantage of petroleum jelly-based products relates to the greasy feeling left following their topical application onto

the skin, mucosal membranes and wounds. Besides petroleum jelly, hydrophobic pharmaceutical carriers now in use include liquid paraffin, lanolin, beeswax, vegetable oil, glycerin monostearate, higher alcohols, polyethylene glycol and some emulsifying agents, which also have undesirable flow properties and skin 5 feel.

Several hydrophobic liquid and semi-solid oils, e.g., mono- and poly-unsaturated oils from vegetable and marine sources, mineral oils, silicone oils, and liquid hydrophobic plant-derived oils, are known for their therapeutic benefits when applied topically, yet, their application in liquid form is not practical. Oils 10 can also contain essential nutritional constituents, such as oil-soluble vitamins (e.g., vitamin A and vitamin E), minerals and other therapeutically beneficial constituents. Another class of therapeutic oils includes mineral and silicon oils useful for the treatment of skin dehydration and other medical disorders, which oils are liquid at ambient temperature. Such therapeutic oils unfortunately, cannot 15 be applied by users in amounts sufficient to exert therapeutic affects because of they typically are liquid at use temperatures.

Other pharmaceutical active ingredients are water-soluble and require a water component in the carrier.

While semi-solid cosmetic and pharmaceutical formulations, such as 20 creams, lotions, gels and ointments are commonly used by consumers, new forms are desirable, in order to achieve better control of the application, while maintaining or bestowing the skin beneficial properties of such products. Thus, the development of a new composition, having breakable foam consistency when extruded out of a container and liquid properties when applied onto the skin is 25 advantageous. Ideally a foam should contain hydrophobic substances (solvents), which can act as emollients and provide the skin with soothing and nourishing properties. However, such hydrophobic solvents are difficult to formulate into a lather-producing or foam-producing product because the hydrophobic solvents interfere with the lather forming ability of the surfactant. Furthermore, addition of 30 oils and other emollients to topical formulations can result in an unpleasant or annoying skin residue.

Use of emulsions in foam compositions is known. Emulsion systems provide a two-phase system including lipophilic or hydrophobic components in one phase and hydrophilic components in the second phase. The foamed emulsion typically is an oil-in-water emulsion in which the hydrophobic 5 component is dispersed in the aqueous continuous phase. Surfactants for reducing surface tension and emulsifiers for improving foam stability are included in the foam composition.

Foams and, in particular, foam emulsions are complicated systems which do not form under all circumstances. Slight shifts in foam emulsion composition, 10 such as by the addition of active ingredients, may destabilize the foam. Furthermore, many emulsions do not provide the high foam capacity, foam stability and/or fast-breaking action under stress or temperatures that are desired in a topical foam composition.

A particularly desirable type of oil-containing foam is such wherein all or 15 part of the oil phase comprises silicone oil. Silicone oil is known for its skin protective features and its incorporation in topical products is beneficial. However, it is not obvious to produce silicone oil-based foams, since many silicone oils possess anti-foaming properties.

US Pat. No. 6,126,920 discloses treatment of various skin diseases, and in 20 particular, scalp psoriasis, using a foamable pharmaceutical composition containing a corticosteroid active substance, an aliphatic alcohol, water, a fatty alcohol, a surface-active agent, a propellant and a buffering agent. The foamable composition contains 40-90% w/w composition of an aliphatic alcohol. US Pat. No. 6,126,920 is typical of many compositions that use aliphatic alcohols in the 25 foam composition. The alcohol promotes fast drying and thereby attempts to address the sticky feeling left by many topical formulations after application; however, alcohols, and in particular the methyl, ethyl and isopropyl alcohols preferred in the '920 patent, are defatting agents and may cause skin to become dry and cracked. Hence, the presence of aliphatic alcohol in a therapeutic foam 30 for external topical administration as taught in US Pat. No. 6,126,920 is undesirable.

US Pat. No. 5,536,743 to Borgman describes a buffered non-flowing composition suitable for the treatment of bacterial vaginosis which contains metronidazole. Suitable formulations include oil-in-water emulsions including an internal oil phase of about 10-40 wt% oil and anionic, cationic or nonionic surfactants. Suitable components of the oleaginous phase include long chain alcohols, esters, and acids, vegetable and animal oils and waxes. No other stabilizing agents are disclosed for use in foam aerosol compositions.

EP 0,598,412 describes a composition that is useful for skin protection against drying and harsh environmental substances. The protection is derived from the inclusion of poly(tetrafluoroethylene) (PTFE) in the composition. The composition includes low levels of both hydrophilic emollients and hydrophobic emollients. The compositions include high levels of surfactants, including ionic surfactants, and co-emulsifiers resulting in thick emulsions which are not flowable, and thus providing products which are inefficient foamers (or non-foaming) and too thick for spreading over large skin areas.

US Pat. 6,423,323 describes an aqueous foam emulsion. The composition includes a hydrophobic phase including fatty acids, emulsifiers and co-emulsifiers, and an aqueous phase containing hydrophilic moisturizers and emulsifiers. An optional ingredient according to US Pat. 6,423,323 is one or more refatting substances, in preferable concentrations of 0.5 to 2%, if the product is to be used for normal skin; and 3 to 6% for dry skin. Addition of high levels of co-emulsifiers such as fatty alcohols and fatty acids suggest that the foam is not stable. No other stabilizing agents are disclosed.

US Pat. 5,635,469 describes a foamable cleansing liquid composition comprising about 0.05% to about 10% of an emollient, in addition to cleansing surfactants, humectants and water soluble cationic or nonionic polymers, but no propellants. Low density foams are achieved using a novel non-aerosol foam dispenser. The foaming is achieved by operating a manual pump, which is not convenient for operation. Emollients and humectants are included to improve the level of hydration and/or lipid content of the skin. However, the patent notes that emollients and humectants interfere with the lather forming ability of the surfactant.

US Pat. 6,113,888 teaches a single water phase composition comprising a self-tanning agent, a nitrogen-free polymer, a nitrogen-free surfactant, and water.

US Pat. 5,679,324 to Lisboa pertains to an aerosol foamable fragrance composition, translucent in its pre-dispensed state, which forms a fast breaking

5 foam. Apparently the foam breaks spontaneously upon discharging from an aerosol container (with no need of any rubbing or sheer force application), thus, making it impractical for spreading over a skin surface. The composition contains surfactant, a propellant, a fragrance, a thickener, and a cosmetic vehicle (preferably water) wherein the ratio of the surfactant to propellant is from about 10 1:1 to about 1:10. Emollients including silicone oils, mineral oils and hydrocarbon oils may be included.

US Pat. 6,251,369 discloses foamable dental fluoride compositions containing a water-soluble fluoride component, whereby said compositions include an oil in water emulsion. However, the patent fails to specify the identity 15 or concentration of the oil component of the emulsion; and none of the compositions presented in the examples contain any oil component.

US Pat. 5,961,957 describes a barrier foam composition comprising from 70 to 90% of water, from 7 to 9% of butane, from 2 to 4% of glyceryl monostearate, from 1.5 to 3.50% of dimethicone copolyol (a water-soluble 20 silicone compound), from 1 to 3% of propane, from 0.5 to 2.5% of lanolin, from 0.5 to 2.5% of stearic acid and from 0.05 to 1.05% of at least one of methylchloroisothiazolinone and methylisothiazolinone.

A few dermatological foam products are available on the market.

Olux™ Foam, produced by Connetics, Inc., contains clobetasol 25 propionate. Each gram of Olux™ Foam contains 0.5 mg clobetasol propionate, USP, in a thermolabile foam, which consists of ethanol (60%), purified water, propylene glycol, cetyl alcohol, stearyl alcohol, polysorbate 60, citric acid, and potassium citrate. It is dispensed from an aluminum can pressurized with a hydrocarbon propellant (propane/butane). Luxiq™ is another corticostroid foam 30 medication, containing 1.2 mg betamethasone valerate per gram, in a vehicle, comprising ethanol (60.4%), purified water, propylene glycol, cetyl alcohol, stearyl

alcohol, polysorbate 60, citric acid, and potassium citrate, and pressurized with a hydrocarbon propellant.

Cortifoam, a hydrocortisone acetate rectal foam is produced by Schwartz Pharma GmbH, wherein the hydrocortisone is present at 10% in a foam vehicle.

5 Nonmedicinal ingredients of Cortifoam include cetyl alcohol, ethoxylated stearyl alcohol, methylparaben, polyoxyethylene-10 stearyl ether, propylene glycol, propylparaben, triethanolamine, water, and inert propellants, isobutene, and propane.

Thus, foam compositions for topical treatment, containing higher
10 concentrations of oils, and do not comprise alcohol are still desirable. Foam compositions that are robust and suitable for inclusion of a wide range of active ingredients are desired.

SUMMARY OF THE INVENTION

Despite the commonly known fact that hydrophobic solvents are difficult to formulate into a lather-producing or foam-producing product and that addition of conventional hydrophobic solvents interferes with the lather forming ability of the

5 surfactant, we have surprisingly discovered a series of foamable carrier compositions, which, upon admixing with a liquefied gas propellant in an aerosol container, produces a foamable composition that is suitable for topical administration. Upon discharge from an aerosol container, the composition forms a breakable foam, which is rich and creamy in appearance, and show very fine

10 bubble structure. The foam does not break down immediately upon discharge, however, it collapses to spread easily onto a skin area upon slight rubbing.

In one or more embodiments of the present invention, the alcohol-free cosmetic or pharmaceutical foamable carrier composition includes water, a liquid, non-volatile hydrophobic solvent, a foam adjuvant agent selected from the group

15 consisting of fatty acids and fatty alcohols, a surface-active agent and a water gelling agent. Such foamable carriers, when placed in an aerosol container and combined with a liquefied gas propellant, create an oil in water emulsion, which, upon release from the aerosol container, provides a therapeutically beneficial foam product. The foam retains its structure for a time sufficient for a user to

20 apply and to rub the foam into the skin. The foam has a very low yield strength and, hence, it breaks upon touch and makes rubbing easy and efficient, and its application even.

In one or more embodiments of the present invention, the foamable carrier composition the hydrophobic solvent content is about 2-5% and has a

25 composition as follows:

Class A Composition:

- about 2-5% hydrophobic solvent;
- about 80-98% water;
- about 0.1% to 5% foam adjuvant agent;
- 30 – about 0.1% to 5% surface-active agent; and
- about 0.1% to 5% water gelling agent.

In one or more embodiments of the present invention, the foamable composition the hydrophobic solvent content is about 5-10% and has a composition as follows:

Class B Composition:

5 – about 5-10% hydrophobic solvent;
– about 75-95% water;
– about 0.1% to 5% foam adjuvant agent;
– about 0.1% to 5% surface-active agent; and
– about 0.1% to 5% water gelling agent.

10 In one or more embodiments of the present invention, the foamable composition the hydrophobic solvent content is about 10-20% and has a composition as follows:

Class C Composition:

15 – about 10-20% hydrophobic solvent;
– about 60-90% water;
– about 0.1% to 5% foam adjuvant agent;
– about 0.1% to 5% surface-active agent; and
– about 0.1% to 5% water gelling agent.

20 In one or more embodiments of the present invention, the foamable composition the hydrophobic solvent content is about 20-75% and has a composition as follows:

Class A Composition:

25 – about 20-75% hydrophobic solvent;
– about 25-75% water;
– about 0.1% to 5% foam adjuvant agent;
– about 0.1% to 5% surface-active agent; and
– about 0.1% to 5% water gelling agent.

All % values are provided on a weight (w/w) basis, based on the composition with out propellant (unless otherwise specified)..

30 The cosmetic or pharmaceutical foamable carrier composition is liquid. The foamable of the present invention does not contain short chain aliphatic

alcohols, making it non-irritating and non-drying. Alcohols penetrate the skin's protective barrier and break down the intercellular matrix. In a recent publication by the American Academy of Dermatology (AAD), titled "Facing the Facts about Skin Care Products" it is stated "[i]ndividuals with dry skin should avoid

5 astringents and any product with alcohol because they easily strip away moisture from the skin" (see: www.aad.org/PressReleases/FacingFacts.html). Another AAD publication, titled "Sensitive About Your Skin?", recommends to "[a]void solvents that penetrate the skin including, propylene glycol and ethanol" (see: www.aad.org/PressReleases/sensitive.html).

10 The alcohol-free foam carrier is formulated as an oil-in-water or water-in-oil emulsion, so that it is suitable for inclusion of either water-soluble and oil soluble active agents (or both). The foamable carrier composition of the present invention, when admixed with a propellant substance in an amount of about 5-25% by weight of the total composition in an aerosol container, produces

15 lightweight breakable foam, suitable for facile application onto the skin, and other body areas, which may accept topically-applied products. Since the propellant, in the pressurized container is in liquid state, upon admixing the foamable carrier composition with the propellant, a stable emulsion, comprising the oil and the propellant (jointly as the "oil phase" component of such emulsion) is formed.

20 In one or more embodiments of the present invention, an alcohol-free cosmetic or pharmaceutical product is provided. The product includes a foam carrier composition according to one or more embodiments of the present invention and an active cosmetic or pharmaceutical ingredient in a therapeutically effective concentration. Cosmetic and pharmaceutical agents can be included in

25 each of the compositions described above and in the detailed description that follows. Pharmaceutical products are intended for topical treatment of human and animal skin disorders, or any other disorder, that requires topical application of a drug. Cosmetic products are intended for beautifying the skin and improving its appearance.

30 Cosmetic and medical disorders that are best treated using the alcohol-free foam carrier and the alcohol-free cosmetic or pharmaceutical product are

identified, and the advantages of such carrier and products is demonstrated as compared to currently available options.

The foam of the present invention is advantageous to current options, for one or more of the following reasons:

- 5 (1) The foam is lightweight and thus, economical.
- (2) The foam contains a hydrophobic solvent, in any desirable concentration, which provides a refatting and skin soothing effect, as well as a carrier for hydrophobic active agents.
- 10 (3) The foam contains silicone oil in a therapeutically effective concentration
- (4) The foam includes active agent, both water soluble and oil soluble.
- (5) The foam is easily spreadable, allowing treatment of large areas such as the arms, back, legs and the breast.
- 15 (6) Due to its flow properties, it spreads effectively into folds and wrinkles, providing uniform distribution of the active agent without the need of extensive rubbing and absorbs into the skin.

BRIEF DESCRIPTION OF THE DRAWINGS

A more complete appreciation of the present invention and many of its advantages will be understood by reference to the following detailed description when considered in connection with the following drawings, which are presented

5 for the purpose of illustration only are not intended to limit the scope of the appended claims, and in which:

Figure 1 illustrates the improvement in the treatment of psoriasis using Bethasone valerate 0.12% foam; and

Figure 2 illustrates the improvement in the treatment of atopic dermatitis

10 using Bethasone valerate 0.12% foam.

DETAILED DESCRIPTION OF THE INVENTION

Hydrophobic solvent

A hydrophobic solvent according to the present invention is a liquid material having solubility in distilled water at ambient temperature of less than

5 about 1 gm per 100 mL, more preferable less than about 0.5 gm per 100 mL, and most preferably less than about 0.1 gm per 100 mL. It is liquid at ambient temperature.

The total content of hydrophobic solvent may vary from 2% to 75% (w/w) of the foamable composition. However, different ranges (herein "composition 10 classes A-D") have been designated, in order to facilitate a choice of an appropriate class, according to the anticipated cosmetic or pharmaceutical need. As a rule of thumb, higher hydrophobic solvent concentrations are more appropriate for the treatment of dry skin, and/or for the treatment of a disease, which is more responsive to drugs delivered in an oily vehicle. Likewise, the 15 higher oil-content composition classes provide an enhanced occlusive effect, which in turn induces the skin penetration of an active agent. Another consideration relates to user acceptance of a product containing a high concentration of the hydrophobic solvent (from about 25% of the composition), which would leave some oily feeling post-application. Thus, a particular 20 composition of the present invention is selected having a hydrophobic solvent concentration in view of the target population and its specific needs.

In one or more embodiments of the present invention, the hydrophobic solvent is mineral oil. Mineral oil (Chemical Abstracts Service Registry number 8012-95-1) is a mixture of aliphatic, naphthalenic, and aromatic liquid 25 hydrocarbons that are derived from petroleum. It is typically liquid; its viscosity is in the range of about 35 CST to about 100 CST (at 40°C), and its pour point (the lowest temperature at which an oil can be handled without excessive amounts of wax crystals forming) is below 0°C. By contrast, white petrolatum, also termed "Vaseline", is disadvantageous, due to its waxy nature. It is known to leave waxy 30 and sticky feeling after application and occasionally stain cloths. Thus, white petrolatum and other semi-solid oils are not a preferred hydrophobic solvent according to the present invention.

Yet another preferred hydrophobic solvents are liquid oils from vegetable, marine or animal sources. By way of example, the unsaturated oil may be selected from the group consisting of olive, corn, soybean, canola, cottonseed, coconut, sesame, sunflower, borage seed, *syzygium aromaticum*, hempseed, 5 herring, cod-liver, salmon, flaxseed, wheat germ and evening primrose oils and mixtures thereof, at any proportion.

A particularly preferred class of oils includes polyunsaturated oils, e.g., esters, and in particular glyceryl esters, of omega-3 and omega-6 fatty acids. Examples of such polyunsaturated fatty acids are linoleic and linolenic acid,

10 gamma-linoleic acid (GLA), eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA). Thus, in one or more embodiments of the present invention the hydrophobic solvent includes at least 6% by weight foamable composition of an oil selected from omega-3 oil, omega-6 oil, and mixtures thereof.

Another class of oils suitable for use as a hydrophobic solvent is liquid 15 hydrophobic plant-derived oils, or essential oils, e.g. "therapeutic oils" containing active biologically occurring molecules that have a therapeutic effect when applied topically. Examples of such oils include rosehip oil, which contain retinoids and is known to reduce acne and post-acne scars, and tea tree oil, which possess antibacterial, antifungal and antiviral properties. Other examples 20 of essential oils are oils of basil, camphor, cardamom, carrot, citronella, clary sage, clove, cypress, frankincense, ginger, grapefruit, hyssop, jasmine, lavender, lemon, mandarin, marjoram, myrrh, neroli, nutmeg, petitgrain, sage, tangerine, vanilla, verbena, as well as any other therapeutically beneficial oil, known in the art of herbal medication.

25 In one or more embodiments of the present invention, the hydrophobic solvent is an "emollient". An emollient is a hydrophobic agent that softens, smoothens and improves lipid content of the skin or other mucous membranes. In one or more embodiments of the present invention, the emollient is a liquid. Without derogating the generality of this definition, examples of suitable 30 emollients for use include isostearic acid derivatives, isopropyl palmitate, lanolin oil, diisopropyl dimerate, diisopropyl adipate, dimethyl isosorbide, maleated soybean oil, octyl palmitate, isopropyl isostearate, cetyl lactate, cetyl ricinoleate,

tocopheryl acetate, acetylated lanolin alcohol, cetyl acetate, phenyl trimethicone, glyceryl oleate, tocopheryl linoleate, wheat germ glycerides, arachidyl propionate, myristyl lactate, decyl oleate, propylene glycol ricinoleate, isopropyl lanolate, pentaerythrityl tetrastearate, neopentylglycol dicaprylate/dicaprate, hydrogenated 5 coco-glycerides, isononyl isononanoate, isotridecyl isononanoate, myristyl myristate, triisocetyl citrate, octyl dodecanol, octyl hydroxystearate and mixtures thereof. Other examples of other suitable emollients can also be found in the Cosmetic Bench Reference, pp. 1.19-1.22 (1996). In one or more embodiments, the hydrophobic solvent is a mixture of a mineral oil or silicone oil and an 10 emollient.

In one or more embodiments of the present invention, silicone oil is a component of the hydrophobic solvent. Silicone oils are used in the foamable compositions due to their known skin protective and occlusive properties. Suitable silicone oils for use in the invention include non-volatile silicones, such 15 as polyalkyl siloxanes, polyaryl siloxanes, polyalkylaryl siloxanes and polyether siloxane copolymers, polydimethylsiloxanes (dimethicones) and poly(dimethylsiloxane)-(diphenyl-siloxane) copolymers. These are preferably chosen from cyclic or linear polydimethylsiloxanes containing from about 3 to about 9, preferably from about 4 to about 5, silicon atoms. Volatile silicones such 20 as cyclomethicones can also be used. Water-soluble silicones, such as dimethicone copolyol are not included in the definition of silicone oils (as hydrophobic solvents) according to the present invention.

In one or more embodiments of the present invention, the composition comprises at least 2% (w/w foamable composition) silicone oil, alone or as part of 25 the hydrophobic solvent. Yet, in other embodiments, the composition comprises at least 5% (w/w) silicone oil alone or as part of the hydrophobic solvent.

The hydrophobic solvent of the present invention may comprise a mixture of two or more of the above hydrophobic solvents in any proportion.

30 **Foam adjuvant agents**

Foam adjuvants are included in the foamable compositions of the present invention to increase the foaming capacity of surfactants and/or to stabilize the

foam. In one or more embodiments of the present invention, the foam adjuvant agents includes fatty alcohols having 15 or more carbons in their carbon chain, such as cetyl alcohol and stearyl alcohol (or mixtures thereof). Other examples of fatty alcohols are arachidyl alcohol (C20), behenyl alcohol (C22), 1-triacontanol (C30), as well as alcohols with longer carbon chains (up to C50). Fatty alcohols, derived from beeswax, including a mixture of alcohols, a majority of which has at least 20 carbon atoms in their carbon chain, are especially well suited as foam adjuvant agents according to the present invention. The concentration of the fatty alcohol, required to support the foam system is inversely related to the 10 length of its carbon chains.

In one or more embodiments of the present invention, the foam adjuvant agent includes fatty acids having 16 or more carbons in their carbon chain, such as hexadecanoic acid (C16) stearic acid (C18), arachidic acid (C20), behenic acid (C22), octacosanoic acid (C28), as well as fatty acids with longer carbon 15 chains (up to C50), or mixtures thereof.

Optionally, the carbon atom chain of the fatty alcohol or the fatty acid may have at least one double bond. A further class of foam adjuvant agent according to the present invention comprises a long chain fatty alcohol or fatty acid, wherein the carbon atom chain is branched. The carbon chain of the fatty acid or fatty 20 alcohol can be substituted with a hydroxyl group, such as 12-hydroxy stearic acid.

The foam adjuvant agent according to one or more embodiments of the present invention includes a mixture of fatty alcohols, fatty acids and hydroxy fatty acids and derivatives thereof in any proportion, providing that the total 25 amount is 0.1% to 5% (w/w) of the carrier mass. More preferably, the total amount is 0.4% - 2.5% (w/w) of the carrier mass.

While fatty alcohols and fatty acids serve to stabilize the resultant foam composition, they often provide additional therapeutic properties. Long chain saturated and mono unsaturated fatty alcohols, e.g., stearyl alcohol, erycycl 30 alcohol, arachidyl alcohol and docosanol have been reported to possess antiviral, anti infective, anti-proliferative and anti-inflammatory properties (US Patent No. 4,874,794). Longer chain fatty alcohols, e.g., tetracosanol, hexacosanol,

heptacosanol, octacosanol, triacontanol, etc. are also known for their metabolism modifying properties and tissue energizing properties. Long chain fatty acids have also been reported to possess anti-infective characteristics. Thus, the pharmaceutical or cosmetic carrier, containing the foam adjuvant agent of the 5 present invention provides an extra therapeutic benefit in comparison with currently used vehicles, which are inert and non-active.

Surface-active agents

Surface-active agents, according to the present invention include any 10 agent linking oil and water in the composition.

The surface-active agent is suitably selected from anionic, cationic, nonionic, zwitterionic, amphoteric and ampholytic surfactants, as well as mixtures of these surfactants. Such surfactants are well known to those skilled in the pharmaceutical and cosmetic formulation art. Nonlimiting examples of possible 15 surfactants include polysorbates, such as polyoxyethylene (20) sorbitan monostearate (Tween 60) and poly(oxyethylene) (20) sorbitan monoleate (Tween 80); poly(oxyethylene) (POE) fatty acid esters, such as Myrij 45, Myrij 49 and Myrij 59; poly(oxyethylene) alkyl ethers, such as poly(oxyethylene) cetyl ether, poly(oxyethylene) palmityl ether, polyethylene oxide hexadecyl ether, 20 polyethylene glycol cetyl ether, brij 38, brij 52, brij 56 and brij W1; sucrose esters, partial esters of sorbitol and its anhydrides, such as sorbitan monolaurate and sorbitan monolaurate; mono or diglycerides, isoceteth-20, sodium methyl cocoyl taurate, sodium methyl oleoyl taurate, sodium lauryl sulfate, triethanolamine lauryl sulfate and betaines.

25 A combination of surface active agents is possible. Any surface-active agent or combinations thereof may be used as surface-active agent. According to one or more embodiments of the present invention, the surface-active agent (or agents) has an HLB of higher than 9.

In one or more embodiments of the present invention, the surface-active 30 agent is selected from the groups of non ionic surfactants, cationic surfactants, amphoteric and zwitterionic surfactants, and, in particular, the surface-active agent is a non-ionic surfactant. Ionic surfactants (including cationic, anionic,

amphoteric and zwitterionic surfactants) are known to be skin irritants. Therefore, non-ionic surfactants are preferred in applications including sensitive skin such as found in most dermatological disorders. We have surprisingly found that non-ionic surfactants alone provide foams of excellent quality, i.e. a score of 5 "E" according to the grading scale discussed below.

In one or more embodiments of the present invention, the surface active agent is solely non-ionic, comprising one or more non-ionic surfactants.

In one or more embodiments of the present invention, the surface active agent include a ratio of non-ionic surfactants to ionic surfactants in the range of

10 100:1 to 6:1; in some embodiments the non-ionic to ionic surfactant ratio is greater than 6:1, or greater than 8:1; or greater than 14:1, or greater than 16:1, or greater than 20:1.

Exemplary non-ionic surfactants include polyethoxylated fatty acids, fatty acid diesters, polyethylene glycol glycerol fatty acid esters, alcohol-oil transesterification products, polyglycerized fatty acids, sterol and sterol derivatives, polyethylene glycol sorbitan fatty acid esters, polyethylene glycol alkyl ethers, sugar esters, polyethylene glycol alkyl phenols, polyoxyethylene-polyoxypropylene block copolymers, sorbitan fatty acid esters and lower alcohol fatty acid esters.

20 Although polyethylene glycol (PEG) itself does not function as a surfactant, a variety of PEG-fatty acid esters have useful surfactant properties. Exemplary monoesters include esters of lauric acid, oleic acid, and stearic acid, e.g., PEG-8 laurate, PEG-8 oleate, PEG-8 stearate, PEG-9 oleate, PEG-10 laurate, PEG-10 oleate, PEG-12 laurate, PEG-12 oleate, PEG-15 oleate, PEG-20 laurate and 25 PEG-20 oleate. Polyethylene glycol fatty acid diesters suitable for use as non-ionic surfactants in the compositions of the present invention include PEG-20 dilaurate, PEG-20 dioleate, PEG-20 distearate, PEG-32 dilaurate and PEG-32 dioleate. Suitable polyethylene glycol glycerol fatty acid esters include PEG-20 glyceryl laurate, PEG-30 glyceryl laurate, PEG-40 glyceryl laurate, PEG-20 30 glyceryl oleate, and PEG-30 glyceryl oleate.

A large number of surfactants of different degrees of hydrophobicity or hydrophilicity can be prepared by reaction of alcohols or polyalcohols with a

variety of natural and/or hydrogenated oils. Most commonly, the oils used are castor oil or hydrogenated castor oil, or an edible vegetable oil such as corn oil, olive oil, peanut oil, palm kernel oil, apricot kernel oil, or almond oil. Preferred alcohols include glycerol, propylene glycol, ethylene glycol, polyethylene glycol, 5 sorbitol, and pentaerythritol. Among these alcohol-oil transesterified surfactants, preferred hydrophilic surfactants are PEG-35 castor oil (Incrocas-35), PEG-40 hydrogenated castor oil (Cremophor RH 40), PEG-25 trioleate (TAGAT® TO), PEG-60 corn glycerides (Crovil M70), PEG-60 almond oil (Crovil A70), PEG-40 palm kernel oil (Crovil PK70), PEG-50 castor oil (Emalex C-50), PEG-50 10 hydrogenated castor oil (Emalex HC-50), PEG-8 caprylic/capric glycerides (Labrasol), and PEG-6 caprylic/capric glycerides (Softigen 767). Preferred hydrophobic surfactants in this class include PEG-5 hydrogenated castor oil, PEG-7 hydrogenated castor oil, PEG-9 hydrogenated castor oil, PEG-6 corn oil (Labrafil® M 2125 CS), PEG-6 almond oil (Labrafil® M 1966 CS), PEG-6 apricot 15 kernel oil (Labrafil® M 1944 CS), PEG-6 olive oil (Labrafil® M 1980 CS), PEG-6 peanut oil (Labrafil® M 1969 CS), PEG-6 hydrogenated palm kernel oil (Labrafil® M 2130 BS), PEG-6 palm kernel oil (Labrafil® M 2130 CS), PEG-6 triolein (Labrafil® b M 2735 CS), PEG-8 corn oil (Labrafil® WL 2609 BS), PEG-20 corn glycerides (Crovil M40), and PEG-20 almond glycerides (Crovil A40). The latter 20 two surfactants are reported to have HLB values of 10, which is generally considered to be the approximate border line between hydrophilic and hydrophobic surfactants.

Alcohol-oil transesterification derivatives of oil soluble vitamins (e.g., vitamins A, D, E, K, etc.), such as tocopheryl PEG-100 succinate (TPGS, 25 available from Eastman), are also suitable surfactants.

Polyglycerol esters of fatty acids are also suitable non-ionic surfactants for the present invention. Among the polyglyceryl fatty acid esters, exemplary use hydrophobic surfactants include polyglyceryl oleate (Plurol Oleique), polyglyceryl-2 dioleate (Nikkol DGDO), and polyglyceryl-10 trioleate. Preferred hydrophilic 30 surfactants include polyglyceryl-10 laurate (Nikkol Decaglyn 1-L), polyglyceryl-10 oleate (Nikkol Decaglyn 1-O), and polyglyceryl-10 mono, dioleate (Caprol® PEG

860), Polyglyceryl polyricinoleates (Polymuls) are hydrophilic and hydrophobic surfactants of this class.

Sterols and derivatives of sterols are suitable surfactants for use in the present invention. These surfactants can be hydrophilic or hydrophobic.

5 Preferred derivatives include the polyethylene glycol derivatives. An exemplary hydrophobic surfactant in this class is cholesterol. An exemplary hydrophilic surfactant in this class is PEG-24 cholesterol ether (Solulan C-24).

A variety of PEG-sorbitan fatty acid esters are suitable for use as non-ionic surfactants in the present invention. In general, these surfactants are hydrophilic, 10 although several hydrophobic surfactants of this class can be used. Among the PEG-sorbitan fatty acid esters, exemplary hydrophilic surfactants include PEG-20 sorbitan monolaurate (Tween-20), PEG-20 sorbitan monopalmitate (Tween-40), PEG-20 sorbitan monostearate (Tween-60), and PEG-20 sorbitan monooleate (Tween-80).

15 Ethers of polyethylene glycol and alkyl alcohols are suitable non-ionic surfactants for use in the present invention. Exemplary hydrophobic ethers include PEG-3 oleyl ether (Volpo 3) and PEG-4 lauryl ether (Brij 30).

The polyoxyethylene-polyoxypropylene (POE-POP) block copolymers are a unique class of polymeric surfactants. The unique structure of the surfactants, 20 with hydrophilic POE and hydrophobic POP moieties in well-defined ratios and positions, provides a wide variety of surfactants suitable for use in the present invention. These surfactants are available under various trade names, including Synperonic PE series (ICI), Pluronic® series (BASF), Emkalyx, Lutrol (BASF), Supronic, Monolan, Pluracare, and Plurodac. The generic term for these 25 polymers is “poloxamer” (CAS 9003-11-6). Exemplary hydrophilic surfactants of this class include Poloxamers 108, 188, 217, 238, 288, 338, and 407. Exemplary hydrophobic surfactants in this class include Poloxamers 124, 182, 183, 212, 331, and 335.

Sorbitan esters of fatty acids are suitable non-ionic surfactants for use in 30 the present invention. Among these esters, preferred hydrophobic surfactants include sorbitan monolaurate (Arlacel 20), sorbitan monopalmitate (Span-40), sorbitan monooleate (Span-80), sorbitan monostearate, and sorbitan tristearate.

Esters of lower alcohols (C₂ to C₄) and fatty acids (C₈ to C₁₈) are suitable non-ionic surfactants for use in the present invention. Among these esters, preferred hydrophobic surfactants include ethyl oleate (Crodamol EO), isopropyl myristate (Crodamol IPM), and isopropyl palmitate (Crodamol IPP).

5 In one or more embodiments of the present invention, the surface-active agent comprise mono-, di- and tri-esters of sucrose with food fatty acids (sucrose esters), prepared from sucrose and methyl and ethyl esters of food fatty acids or by extraction from sucroglycerides. Exemplary sucrose esters include sucrose monopalmitate and sucrose monolaurate. Suitable sucrose esters include those
10 having a high monoester content, which have higher HLB values.

In one or more embodiments of the present invention, a combination of a non-ionic surfactant and an anionic surfactant (such as sodium lauryl sulphate) is employed, at a ratio of between 1:1 and 20:1, or at a ratio of 4:1 to 10:1. The resultant foam has a low specific gravity, e.g., less than 0.1g/ml, which upon
15 rubbing (shear stress) onto the skin collapses easily, to allow facile absorption.

Unlike prior art foamable compositions, the total surfactant employed to obtain a foam that is stable, of low specific gravity and has a fine bubble structure is low. Lower surfactant levels, particularly of ionic surfactants, are preferred to reduce skin irritations. Total surfactant is in the range of 0.1 to 5.0 wt% of the
20 foamable composition, and is typically less than 2 wt%, or even less than 1 wt%.

Water gelling agents

The water gelling agent according to one or more embodiments of the present invention stabilizes the aqueous phase by, for example, increasing
25 viscosity and linking capability. Exemplary water gelling agents that can be used in accordance with one or more embodiments of the present invention include for example, but are not limited to, naturally-occurring polymeric materials such as, locust bean gum, sodium alginate, sodium caseinate, egg albumin, gelatin agar, carrageenin gum sodium alginate, xanthan gum, quince seed extract, tragacanth
30 gum, starch, chemically modified starches and the like, semi-synthetic polymeric materials such as cellulose ethers (e.g. hydroxyethyl cellulose, methyl cellulose, carboxymethyl cellulose, hydroxy propylmethyl cellulose), polyvinylpyrrolidone,

polyvinylalcohol, guar gum, hydroxypropyl guar gum, soluble starch, cationic celluloses, cationic guar and the like and synthetic polymeric materials such as carboxyvinyl polymers, polyvinylpyrrolidone, polyvinyl alcohol polyacrylic acid polymers, polymethacrylic acid polymers, polyvinyl acetate polymers, polyvinyl chloride polymers, polyvinylidene chloride polymers and the like. Mixtures of the above compounds are contemplated.

5 Further exemplary water gelling agents include the acrylic acid/ethyl acrylate copolymers and the carboxyvinyl polymers sold, for example, by the B.F. Goodrich Company under the trademark of Carbopol Registered TM resins.

10 These resins consist essentially of a colloidal water-soluble polyalkenyl polyether crosslinked polymer of acrylic acid crosslinked with from 0.75% to 2% of a crosslinking agent such as polyallyl sucrose or polyallyl pentaerythritol. Examples include Carbopol 934, Carbopol 940, Carbopol 950, Carbopol 980, Carbopol 951 and Carbopol 981. Carbopol 934 is a water-soluble polymer of acrylic acid

15 crosslinked with about 1% of a polyallyl ether of sucrose having an average of about 5.8 allyl groups for each sucrose molecule.

20 The gelling agent is present in an amount in the range of about 0.1% to about 5.0 wt% of the foammable composition. In one or more embodiments, it is typically less than 1 wt% of the foammable composition.

25 **“Alcohol free”**

Unlike the composition disclosed in US Pat. No. 6,126,920, which contains a 40-90 wt% aliphatic alcohol, the composition of the present invention does not contain such amount alcohols. For the purpose of the present application, the term “alcohol free” shall mean that the composition contains no more than an incidental amount of an aliphatic alcohol, e.g. less than about 7.5% of any aliphatic alcohol, having one to six carbon atoms in their carbon backbone, or no more than 7.5% of any mixture of such aliphatic alcohols. Alcohols at these low levels are not considered to have a negative effect on skin or mucous membranes. In one or more embodiments, the foammable compositions do not contain any alcohol.

Optional Ingredients

The pharmaceutical or cosmetic foam carrier of the present invention may further optionally comprise a variety of pharmaceutical or cosmetic ingredients, which are added in order to fine-tune the consistency of the formulation, protect 5 the formulation components from degradation and oxidation and bestow their cosmetic acceptability. Such excipients, may be selected, for example, from the group consisting of diglycerides, triglycerides, stabilizing agents, antioxidants, humectants, flavoring, colorant and odorant agents and other formulation components, used in the art of pharmaceutical and cosmetic formulary. A 10 pharmaceutical or cosmetic composition manufactured using the foam carrier according to the present invention is very easy to use. When applied onto the afflicted body surface of humans or animals, it is in a foam state, allowing free application without spillage. Upon further application of a mechanical force, e.g., by rubbing the composition onto the body surface, it freely spreads on the surface 15 and is rapidly absorbed.

Propellant Aerosol

Aerosol propellants are used to generate and administer the foamable composition as a foam. The total composition including propellant, foamable 20 compositions and optional ingredients is referred to as the foamable carrier. The propellant makes up about 5-25 wt% of the foamable carrier. Examples of suitable propellants include volatile hydrocarbons such as butane, propane, isobutane or mixtures thereof, and fluorocarbon gases.

25 Composition and Foam Physical Characteristics

1. Composition flow properties:

It is important to have a composition, including water, hydrophobic solvents, formulation excipients and propellant, in a stable emulsion, which ascertain acceptable shelf-life of the product.

30 Yet, another crucial property is that said composition has to be free flowing, since otherwise, it cannot flow through the dip-tube of the aerosol container and create acceptable foam. It has been noted that in the context of the

composition of the present invention, compositions comprising semi-solid hydrophobic solvents, e.g., white petrolatum, are excessively viscous and demonstrate poor flowability.

The combination of a surface active agent, foaming adjuvant and water

5 gelling agent according to one or more embodiments of the invention provides a low specific gravity foam having superior flow properties and sheer breakability (among other attributes). According to one or more embodiments of the present invention, the total amount of surface active agent, foaming adjuvant and water gelling agent, in combination does not exceed 8 % (w/w) of foamable

10 composition. In other embodiments, the combined amounts of surface active agent, foaming adjuvant and water gelling agent is less than 5 % (w/w) of foamable composition. The low solids content improves the flow properties of the foam, reduces unpleasant skin residue and reduces the cost of manufacture. As is demonstrated herein, the foam quality and foam breakability is excellent,

15 despite the low levels of these components in the foam.

2. Foam properties:

The following scale for foam quality is used to evaluate foams.

E (excellent): very rich and creamy in appearance, does not show any bubble structure or shows a very fine (small) bubble structure.

20 **G** (good): rich and creamy in appearance, very small bubble size, "dulls" more rapidly than an excellent foam.

FG (fairly good): a moderate amount of creaminess noticeable, bubble structure is noticeable.

F (fair): very little creaminess noticeable, larger bubble structure than a

25 "fairly good" foam.

P (poor): no creaminess noticeable, large bubble structure.

VP (very poor): dry foam, large very dull bubbles, difficult to spread on the skin.

Foams, adequate for topical administration according to the present

30 invention have to be of quality grade E or G, upon release from the aerosol container. Smaller bubbles mean more stable foam, which does not collapse

spontaneously immediately upon discharge from the container. The finer foam structure looks and feels smoother, thus increasing its usability and appeal.

A crucial aspect of foam properties, according to the present invention is breakability. Sheer-force breakability of the foam, as attained by the composition 5 of the present invention is clearly advantageous to thermally-induced breakability, present, for example in US Pat. 6,126,920, and the respective Olux and Luxiq products, as demonstrated by the fact that according to the use instructions of Olux and Luxiq, the foam cannot be applied on the hand and afterwards delivered to the afflicted area, since it immediately collapses upon exposure to skin 10 temperature.

Yet, another important property is specific gravity of the foam, as measured upon release from the aerosol can. Typically, foams according to the present invention have specific gravity of less than 0.1 g/mL and more preferably, less than 0.05 g/mL.

15

FIELDS OF PHARMACEUTICAL APPLICATIONS

By including an appropriate therapeutic agent in the foamable carrier, the foam composition of the present invention is useful in the therapy of a variety of dermatological disorders (also termed "dermatoses"), including, in a non-limiting 20 exemplary manner:

Dermatitis

- Contact Dermatitis
- Atopic Dermatitis
- Seborrheic Dermatitis
- Nummular Dermatitis
- Chronic Dermatitis Of The Hands And Feet
- Generalized Exfoliative Dermatitis
- Stasis Dermatitis
- Lichen Simplex Chronicus

25

Bacterial Infections

- Cellulitis

- Acute Lymphangitis
- Lymphadenitis
- Erysipelas
- Cutaneous Abscesses
- 5 – Necrotizing Subcutaneous Infections
- Staphylococcal Scalded Skin Syndrome
- Folliculitis
- Furuncles
- Hidradenitis Suppurativa
- 10 – Carbuncles
- Paronychial Infections
- Erythrasma

Fungal Infections

- Dermatophyte Infections
- 15 – Yeast Infections

Parasitic Infections

- Scabies
- Pediculosis
- Creeping Eruption

20 Viral Infections**Disorders of Hair Follicles and Sebaceous Glands**

- Acne
- Rosacea
- Perioral Dermatitis
- 25 – Hypertrichosis (Hirsutism)
- Alopecia, including male pattern baldness, alopecia areata, alopecia universalis and alopecia totalis
- Pseudofolliculitis Barbae
- Keratinous Cyst

30 Scaling Papular Diseases

- Psoriasis

- Pityriasis Rosea
- Lichen Planus
- Pityriasis Rubra Pilaris

Benign Tumors

- 5 – Moles
- Dysplastic Nevi
- Skin Tags
- Lipomas
- Angiomas
- 10 – Pyogenic Granuloma
- Seborrheic Keratoses
- Dermatofibroma
- Keratoacanthoma
- Keloid

15 Malignant Tumors

- Basal Cell Carcinoma
- Squamous Cell Carcinoma
- Malignant Melanoma
- Paget's Disease Of The Nipples
- 20 – Kaposi's Sarcoma

Reactions To Sunlight

- Sunburn
- Chronic Effects of Sunlight
- Photosensitivity

25 Bullous Diseases

- Pemphigus
- Bullous Pemphigoid
- Dermatitis Herpetiformis
- Linear Immunoglobulin A Disease

30 Pigmentation Disorders

- Hypopigmentation

- Vitiligo
- Albinism
- Postinflammatory hypopigmentation
- Hyperpigmentation

5

- Melasma (chloasma)
- Drug-induced hyperpigmentation
- Postinflammatory hyperpigmentation

Disorders of Cornification

- Ichthyosis
- Keratosis Pilaris
- Calluses And Corns
- Actinic keratosis

Pressure Sores**Disorders of Sweating****15 Inflammatory reactions**

- Drug Eruptions
- Toxic Epidermal Necrolysis
- Erythema Multiforme
- Erythema Nodosum

20 – Granuloma Annulare

In one or more embodiments of the present invention, the foam composition of the present invention is useful in the therapy of non-dermatological disorders, which respond to transdermal delivery of an active agent. By way of example, such disorders include localized pain in general, as well as joint pain, muscle pain, back pain, rheumatic pain, arthritis, osteoarthritis and acute soft tissue injuries and sports injuries. Other disorders of this class include conditions, which respond to hormone therapy, such as hormone replacement therapy, transdermal nicotine administration, and other respective disorders, known in the art of drug delivery. The foam composition of the present invention is also useful in the delivery of local anesthetic agents.

ACTIVE PHARMACEUTICAL AGENTS (DRUGS)

The active pharmaceutical agents, also referred to as "drug(s)", may consist of a single drug or a combination of drugs that can be dissolved in the water phase or the hydrophobic phase of the carrier composition. Examples of 5 such drugs are antibiotic, antibacterial, antifungal, antiviral, antiinflammatory, anesthetic, analgesic, antiallergic, corticosteroid, retinoid and antiproliferative medications and mixtures thereof at any proportion. The concentration of drugs may be adopted to exert a therapeutic effect on a disease when applied to an afflicted area.

10 **Antibacterial agents**

One important class of drugs comprises antibacterial agents. It is well known that bacterial infections are involved in a variety of superficial disorders of the skin, eye, mucosal membrane, oral cavity, vagina and rectum.

15 The antibacterial drug can be active against gram positive and gram-negative bacteria, protozoa, aerobic bacteria and unaerobic ones.

By way of example, the antibacterial drugs can be selected from the group of chloramphenicol, tetracyclines, synthetic and semi-synthetic penicillins, beta-lactames, quinolones, fluoroquinolones, macrolide antibiotics, metronidazole and its derivatives and analogs, dicarboxylic acids, such as azelaic acid, siccylates, 20 peptide antibiotics, cyclosporines and any combination thereof at a therapeutically effective concentration. Another group of antibacterial agents which is non-specific, comprises strong oxidants and free radical liberating compounds, such as hydrogen peroxide, bleaching agents (e.g., sodium, calcium or magnesium hypochloride and the like) iodine, chlorohexidine and benzoyl 25 peroxide.

Antibacterial compositions according to the present invention may be used to treat infections of the skin. An example of a very common skin infection is acne, which involve infestation of the sebaceous gland with *p. acnes*, as well 30 *staphylococcus aurus* and *pseudomonas*. Various antibacterial agents have been utilized to treat acne, however, their efficacy is limited due to their low penetration into the hydrophobic environment of the skin layers and sebaceous glands. The composition of the present invention, comprising a hydrophobic component,

would facilitate an enhanced rate of penetration. Furthermore, the intrinsic antibacterial and antiinflammatory effects of the foam adjuvant agents, i.e., fatty alcohols and acids, provides a combined effect that should result in a better therapeutic response to treatment.

5 The composition of the present invention is particularly useful and beneficial in the prevention and treatment of secondary infections, accompanying skin-structure damage, such as in cuts, wounds, burns and ulcers. In all such cases, the present formulation is easy to use, being in foam state when applied and becoming liquid instantly upon rubbing onto the skin.

10 While being useful in the prevention and treatment of infections, the antibacterial foam of the present invention is also applicable for decontaminating areas, afflicted with bacterial warfare organisms, such as anthrax and smallpox.

15 The same advantage is expected when the composition of the present invention is topically applied to mucosal membranes, the oral cavity, the vagina and the rectum.

Anti-fungal agents

20 Fungal infections are another object of treatment using the composition of the present invention. Superficial fungal infection of the skin is one of the commonest skin diseases seen in general practice. Dermatophytosis is probably the most common superficial fungal infection of the skin. It is caused by a group of fungi, which are capable of metabolizing the keratin of human epidermis, nails or hair. There are 3 genera of dermatophytes causing dermatophytosis, i.e., microsporum, trichophyton and epidermophyton.

25 Candidiasis is an infection caused by the yeast like fungus candida albicans or occasionally other species of candida. Clinical syndromes of candidiasis include: (a) oral candidiasis (oral thrush); (b) candidiasis of the skin and genital mucous membrane; and (c) candida paronychia, which inflicts the nail.

30 The pharmaceutical composition may comprise an antifungal drug, which is active against dermatophytes and candida, selected from the group of, but not limited to azoles, diazoles, triazoles, miconazole, fluconazole, ketoconazole, clotrimazole, itraconazole griseofulvin, ciclopirox, amorolfine, terbinafine,

Amphotericin B, potassium iodide, flucytosine (5FC) and any combination thereof at a therapeutically effective concentration.

It is useful, for example for the treatment of tinea corporis, tinea pedis, tinea rubrum, tinea unguium, tinea cruris, tinea barbae and tinea versicolor, as
5 well as yeast infections, such as candidiasis, and candidal vaginitis

Anti-viral agents

The composition of the present invention is particularly beneficial in the case of viral infections. Cold sores are caused by the herpes simplex Type 1 virus and are sometimes referred to as facial herpes. Mollusca are small viral
10 growths that appear singly or in groups on the face, trunk, lower abdomen, pelvis, inner thighs, or penis. Shingles (herpes zoster), which usually only occurs once in a lifetime, appears as a rash (clusters of blisters with a red base). It is caused by the same virus responsible for chickenpox. Warts are a common, benign skin tumor caused by viral infection.

15 Viral infections are currently treated with various antiviral agents, as summarized in the following table:

Drug	Viruses	Chemical Type
Vidarabine	Herpesviruses	Nucleoside analogue
Acyclovir	Herpes simplex (HSV)	Nucleoside analogue
Gancyclovir	Cytomegalovirus (CMV)	Nucleoside analogue
Nucleoside-analog reverse transcriptase inhibitors (NRTI): AZT (Zidovudine), ddI (Didanosine), ddC (Zalcitabine), d4T (Stavudine), 3TC (Lamivudine)	Retroviruses (HIV)	Nucleoside analogue
Non-nucleoside reverse transcriptase inhibitors (NNRTI): Nevirapine, Delavirdine	Retroviruses (HIV)	Nucleoside analogue
Protease Inhibitors: Saquinavir, Ritonavir, Indinavir, Nelfinavir	HIV	Peptide analogue
Ribavirin	Broad spectrum: HCV, HSV, measles, mumps, Lassa fever	Triazole carboxamide
Amantadine / Rimantadine	Influenza A strains	Tricyclic amine
Interferons	Hepatitis B and C	Protein

Any of the above antiviral drugs, in a therapeutically effective concentration, can be incorporated in the foam composition of the present invention. The composition of the present invention, which comprises a

5 hydrophobic solvent, would facilitate an enhanced rate of penetration and better topical distribution of any of the above listed antiviral drugs. Furthermore, the intrinsic antiviral effects of the foam adjuvant agents, i.e., fatty alcohols and acids, provides a combined effect that should result in a better therapeutic response to treatment.

10 **Antiinflammatory or antiallergic agents**

Yet, according to another embodiment according to the present invention the drug is an antiinflammatory or antiallergic agent. Antiinflammatory or antiallergic agent can be selected from the group of corticosteroids, non-steroidal antiinflammatory drugs (NSAIDs), anti-histamines, immunosuppressants and any

15 combination thereof at a therapeutically effective concentration.

The following table provides a summary of currently available corticosteroid agent and their typical therapeutically effective concentration.

Potency	Compound	Formulation
Very high	Clobetasol propionate	Cream or ointment 0.05%
	Halobetasol propionate	Cream or ointment 0.05%
High	Betamethasone dipropionate	Cream or ointment 0.05%
	Betamethasone valerate	Ointment 0.1%
	Fluocinolone acetonide	Cream 0.02%
	Halcinonide	Cream or ointment 0.1%
Medium	Betamethasone valerate	Cream 0.1%
	Fluocinolone acetonide	Cream or ointment 0.020%
	Hydrocortisone valerate	Cream or ointment 0.2%
	Triamcinolone acetonide	Cream, ointment, or lotion 0.1% or 0.020%
Low	Hydrocortisone	Cream, ointment, or lotion 1.0% or 2.5%

5 The concentrations of corticosteroid drugs, as presented in the above table are provided herein only as example, and any therapeutically effective concentration of such corticosteroids can be incorporated in the composition of the present invention.

Since all corticosteroid drugs are typically hydrophobic, the carrier of the
 10 present invention, comprising a hydrophobic solvent, is most suitable as a vehicle to facilitate better topical distribution and an enhanced rate of penetration of any of the above listed drugs. Furthermore, the intrinsic antiviral, antibacterial and antiinflammatory effects of the foam adjuvant agents, i.e., fatty alcohols and acids, provides a combined effect that should result in a better therapeutic
 15 response to treatment.

Psoriasis is a very common chronic skin disease, which may be the target of treatment using the composition of the present invention. It is marked by periodic flare-ups of sharply defined red patches covered by a silvery, flaky surface.

Corticosteroid ointments, greasy preparations containing little or no water, are commonly used for treating psoriasis. Their main disadvantage is in their sticky feeling, which remains so long after treatment is over. By contrast, the foam of the present invention, while comprising considerable concentration of an oil (hydrophobic solvent), spreads very easily throughout the afflicted area and absorbs into the skin without leaving any untoward sensation or look. Examples of other inflammatory disorders, which can be treated by the composition of the present invention, wherein the drug is a steroid are atopic dermatitis, seborrhea, seborrheic dermatitis of the face and trunk, seborrheic blepharitis, contact dermatitis, stasis dermatitis (gravitational eczema; varicose eczema), exfoliative dermatitis (erythroderma), lichen simplex chronicus, pityriasis rosea and pemphigus.

Topical antihistaminic preparations currently available include 1% and 2% diphenhydramine (Benadryl® and Caladryl®), 5% doxepin (Zonalon®) cream, 15 phrilamine maleate, chlorpheniramine and tripelennamine, phenothiazines, promethazine hydrochloride (Phenergan®) and dimethindene maleate. These drugs, as well as additional antihistamines can also be incorporated in the composition of the present invention.

It is pointed out that polyunsaturated fatty acids, containing omega-3 and 20 omega-6 fatty acids (e.g., linoleic and linolenic acid, gamma-linoleic acid (GLA), eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA) are beneficial in the treatment of psoriasis and other skin inflammation conditions.

A second class of anti-inflammatory agents, which is useful in the foam of the present invention, includes the nonsteroidal anti-inflammatory agents 25 (NSAIDs). The variety of compounds encompassed by this group is well-known to those skilled in the art. Specific non-steroidal anti-inflammatory agents useful in the composition invention include, but are not limited to:

- 1) Oxicams, such as piroxicam, isoxicam, tenoxicam, sudoxicam;
- 2) Salicylates, such as salicylic acid, ethyl salicylate, methyl salicylate,

30 aspirin, disalcid, benorylate, trilisate, safapryn, solprin, diflunisal, and fendosal;

3) Acetic acid derivatives, such as diclofenac, fenclofenac, indomethacin, sulindac, tolmetin, isoxepac, furofenac, tiopinac, zidometacin, acematacin, fentiazac, zomepirac, clindanac, oxepinac, felbinac, and ketorolac;

4) Fenamates, such as mefenamic, meclofenamic, flufenamic, niflumic, and tolfenamic acids;

5 5) Propionic acid derivatives, such as ibuprofen, naproxen, benoxaprofen, flurbiprofen, ketoprofen, fenoprofen, fenbufen, indoprofen, pirprofen, carprofen, oxaprozin, pranoprofen, miroprofen, tioxaprofen, suprofen, alminoprofen, and tiaprofenic; and

10 6) Pyrazoles, such as phenylbutazone, oxyphenbutazone, feprazole, azapropazone, and trimethazone.

Any further steroidal and nonsteroidal compounds, having the capacity to prevent, alleviate the symptoms of, treat or cure inflammation processes, are generally included, as possible anti-inflammatory agents, according to the 15 present invention.

The pharmaceutical composition of the present invention may also comprise an antiinflammatory or antiallergic agent, wherein said agent reduces the occurrence of pro-inflammatory cytokines or inhibits the effect of pro-inflammatory cytokines.

20 Mixtures of such anti-inflammatory agents may also be employed, as well as the dermatologically acceptable salts, esters, amides, prodrugs and derivatives of these agents.

Topical application of a foam, comprising a safe and effective dose of an NSAID can be useful in the prevention and/or alleviation of the symptoms of 25 rheumatoid arthritis, osteoarthritis and pain. Topical NSAIDs, incorporated in the foam of the present invention can be also used in the treatment of dermatological disorders, such as acne, rosacea, hair growth disorders, actinic keratosis and certain skin cancer conditions.

Topical Anesthetics

30 The compositions of the present invention may contain a safe and effective amount of a topical anesthetic. Examples of topical anesthetic drugs include benzocaine, lidocaine, bupivacaine, chlorprocaine, dibucaine, etidocaine,

mepivacaine, tetracaine, dyclonine, hexylcaine, procaine, cocaine, ketamine, pramoxine, phenol, and pharmaceutically acceptable salts thereof. Mixtures of such anesthetic agents may be synergistically beneficial.

Keratolytically active agents

5 The term "keratolytically active agent" is used herein to mean a compound which loosens and removes the stratum corneum of the skin, or alters the structure of the keratin layers of skin.

10 Keratolytically active agents are used in the treatment of many dermatological disorders, which involve dry skin, hyperkeratinization (such as psoriasis), skin itching (such as xerosis), acne and rosacea.

15 Suitable keratolytically active agent include phenol and substituted phenolic compounds. Such compounds are known to dissolve and loosen the intracellular matrix of the hyperkeratinized tissue. As such, they are used in the treatment of dermatological disorders. Dihydroxy benzene and derivatives thereof have been recognized as potent keratolytic agents. Resorcinol (m-dihydroxybenzene) and derivatives thereof are used in anti-acne preparations. Hydroquinone (p-dihydroxybenzene), besides its anti-pigmentation properties, is also keratolytic. These compounds also exhibit antiseptic properties. Cresols also possess bactericidal and keratolytic properties.

20 Vitamin A and its derivatives, such as retinoic acid, isoretinoic acid, retinol and retinal are another preferred class of keratolytically active agents.

25 Another group of keratolytically active agents include alpha-hydroxy acids, such as lactic acid and glycolic acid and their respective salts and derivatives; and beta-hydroxy acids, such as Salicylic acid (o-hydroxybenzoic acid) and its salts and pharmaceutically acceptable derivatives, which typically possess anti-inflammatory, as well as keratolytic, activity.

Yet, another class of preferred keratolytically active agents includes urea and its derivatives.

Retinoids

30 Another preferred group of active agents comprise retinol, retinal, all trans retinoic acid and derivatives, isomers and analogs thereof, collectively termed "retinoids". Etretinate, actiretin, isotretinoin, adapalene and tazarotene are further

examples of said retinoid isomers and analogs. Compositions according to the present invention, which contain retinoids as the active drug, can be used for the treatment of acne, seborrhea, various dermatoses, inflammation of the skin, mucosal membranes, vagina and the rectum, psoriasis, actinic keratosis and skin 5 cancers, by application onto the affected area.

Insecticide and Insect repellents agents

Insects, such as mosquitoes, biting flies, mites, gnats, fleas, chiggers, punkies, sand flies, lice and ticks can be annoying and sometimes pose a serious risk to human and animal health. In certain areas of the United States, 10 mosquitoes can transmit diseases like equine and St. Louis encephalitis. Biting flies can inflict a painful bite that can persist for days, swell, and become infected. Ticks can transmit serious diseases like Lyme disease and Rocky Mountain spotted fever.

There are several types of insect repellents to use when protecting people 15 and animals from flying or biting insects, spiders, ticks and mites. By way of example, these may include DEET (N, N-diethyl-m-toluamide), dimethyl phthalate, piperonyl butoxide and permethrin. Insect repelling terpenoids, have been reported by Hwang, et al, J. Chem. Ecol., 11, 1297 (1985); and Ruledge, J. Am. Mosquito Control Assoc. 4, 414 (1988).

20 A particularly preferred group of insect repellents includes the terpenoid compounds, described in U.S. Patent No. 5,411,992, including:

(1) Terpenoid-alcohol or terpene-ols are terpenoids which have at least one hydroxyl group. Examples of terpene-ols include: $C_{10}H_{16}O$ compounds, perillyl alcohol, carveol, myrtenol, and cis-verbenol; $C_{10}H_{18}O$ compounds, 25 myrtanol, iso-pinocampheol, dihydrocarveol, isopulegol, terpineol, terpinen-4-ol, nerol, geraniol, and linalool, and $C_{10}H_{20}O$ compounds, menthol, beta-citronellol, and dihydro-myrcenol.

(2) Terpenoid-esters are terpenoids, which have at least one ester group which is the product of the bonding of the hydroxyl group of a terpene-ol with an 30 aliphatic carboxylic acid that can contain functional groups such as the hydroxyl or amine on the aliphatic chain. Examples of suitable aliphatic carboxylic acids include acetic acid, propionic acid, lactic acid, and various amino acids.

Examples of terpenoid-esters include: carvyl acetate, carvyl propionate, and menthyl lactate.

(3) Essential oils which contain terpenoids and perfumes which contain terpenoids. Non-limiting examples of essential oils which have high content of 5 terpene-ols and esters include bergamot (62% terpenoids); sage (>50% terpenoids); styrax (>50% terpenoids); peppermint (>50% terpenoids); and pine Siberian (75% terpenoids %). Terpenes, aldehydes and ketones vary in their usefulness but as a general group have potential as insect-repellent.

The foam of the present invention is particularly suitable for the effective 10 uniform spreading of an insect repellent agent onto large areas of the skin of humans and animals. The hydrophobic solvent present in the foam composition helps retain the insect repellent on the skin surface for an extended period of time.

Yet, in a further embodiment, the foam is suitable for delivery of insect- 15 killing agents (insecticides) to an afflicted external surface area of humans and animals. Thus, the pharmaceutical or cosmetic composition may comprise an insecticide, known in the art of parasitology. By way of example, such insecticide can be selected selected from the group of permethrin, hexachlorobenzene, carbamate, naturally occurring pyrethroids, permethrin, allethrin, malathion, 20 piperonyl butoxide and any combination thereof at a therapeutically effective concentration. Its application is very convenient and it spreads easily, even over hairy areas. The hydrophobic solvent present in the foam composition helps retain the insecticide on the treated area for an extended period of time. Furthermore, the presence of a hydrophobic solvent in the foam eases 25 mechanical removal of lice and nits with a comb.

Anti cancer drugs

Anti cancer drugs can also be used according to the present invention as the drug of choice from skin malignant tumors, such as basal cell carcinoma, squamous sell carcinoma, melanoma and Kaposi's sarcoma, as well as the pre- 30 cancerous condition actinic keratosis. In certain cases, topical cytotoxic and antiproliferative drugs are used to treat or prevent such cancers, including 5-fluorouracil, also called 5-FU. 5-FU, as well as any other anti-cancer agents,

know in the art of cancer medicine, can be incorporated in the foam at therapeutically effective levels.

A preferred family of anticancer drugs, suitable for usage in the foam of the present formulation comprises antiestrogens, such as tamoxifen. Tamoxifen

5 blocks the effects of the hormone estrogen in the body. It is used to prevent or delay the return of breast cancer or to control its spread.

Photodynamic therapy agents

The foam composition of the present invention is also useful to deliver photo-sensitizing agents, known in the art of photodynamic therapy. By way of

10 example, such photosensitizers can be selected from the group comprising modified porphyrins, chlorins, bacteriochlorins, phthalocyanines, naphthalocyanines, pheophorbides, purpurins, m-THPC, mono-L-aspartyl chlorin e6, bacteriochlorins, phthalocyanines, benzoporphyrin derivatives, as well as photosensitiser precursors, such as aminolevulinic acid (ALA).

15 **Active agents for burns, wounds, cuts and ulcers**

The treatment of burns, wounds, cuts and ulcers, using the composition of the present invention is particularly advantageous. The foam can include both anti-infective agents (against bacteria, fungi and/or viruses), antiinflammatory agents (steroidal and/or NSAIDs) and pain relieving components. Upon

20 application, the foam spreads easily, covering the surface of the affected area, and without causing pain.

SKIN CARE ACTIVE AGENTS

The foam of the present invention is useful and advantageous for skin care

25 and cosmetic care. The combination of oil and water, having moisture-retaining properties, in a spreadable foam form, can be used to substitute currently used cosmetic skin care creams, lotions, gels, etc. The cosmetic foam compositions of the present invention are suitable for the further application as "cosmeceutical" preparation (cosmetic products with therapeutic benefit), to treat "cosmetic" skin

30 disorders, such as aging skin, wrinkles, hyperpigmentation (melasma, chloasma, freckles, etc.), scaly skin and other skin undesirable properties.

The CTFA Cosmetic Ingredient Handbook describes a wide variety of nonlimiting cosmetic and pharmaceutical ingredients commonly used in the skin care industry, which are suitable for use in the compositions of the present invention. Examples of these ingredient classes include: abrasives, absorbents, 5 aesthetic components such as fragrances, pigments, colorings/colorants, essential oils, astringents, etc. (e.g., clove oil, menthol, camphor, eucalyptus oil, eugenol, menthyl lactate, witch hazel distillate), anti-acne agents, antimicrobial agents (e.g., iodopropyl butylcarbamate), antioxidants, binders, biological additives, buffering agents, bulking agents, chelating agents, chemical additives, 10 colorants, cosmetic astringents, cosmetic biocides, denaturants, drug astringents, external analgesics, film formers or materials, e.g., polymers, for aiding the film-forming properties and substantivity of the composition (e.g., copolymer of eicosene and vinyl pyrrolidone), opacifying agents, pH adjusters, propellants, reducing agents, sequestrants, skin bleaching and lightening agents (e.g., 15 hydroquinone, kojic acid, ascorbic acid, magnesium ascorbyl phosphate, ascorbyl glucosamine), skin-conditioning agents (e.g., humectants, including miscellaneous and occlusive), skin soothing and/or healing agents (e.g., panthenol and derivatives (e.g., ethyl panthenol), aloe vera, pantothenic acid and its derivatives, allantoin, bisabolol, and dipotassium glycyrrhizinate), skin treating 20 agents, thickeners, and vitamins and derivatives thereof.

In any embodiment of the present invention, however, the active agents useful herein can be categorized by the benefit they provide or by their postulated mode of action. It is to be understood that the active agents useful herein can in some instances provide more than one benefit or operate via more than one 25 mode of action. Therefore, classifications herein are made for the sake of convenience and are not intended to limit the active to that particular application or applications listed.

Anti-acne active agents

The compositions of the present invention may contain a safe and 30 effective amount of one or more pharmaceutically or cosmetically acceptable anti-acne active agents. Examples of useful anti-acne actives include resorcinol, sulfur, salicylic acid and salicylates, alpha-hydroxy acids, nonsteroidal anti-

inflammatory agents, benzoyl peroxide, retinoic acid, isoretinoic acid and other retinoid compounds, adapalene, tazarotene, azelaic acid and azelaic acid derivatives, antibiotic agents, such as erythromycin and clindamycin, zinc salts and complexes, and combinations thereof, in a therapeutically effective 5 concentration.

Anti-wrinkle active agents/anti-atrophy active agents and agents to treat dry and scaly skin (xerosis and ichthyosis)

The compositions of the present invention may further contain a safe and effective amount of one or more anti-wrinkle actives or anti-atrophy actives, which 10 can be easily delivered by spreading a foam onto the skin. Exemplary anti-wrinkle/anti-atrophy active agents suitable for use in the compositions of the present invention include sulfur-containing D and L amino acids and their derivatives and salts, particularly the N-acetyl derivatives; thiols; hydroxy acids (e.g., alpha-hydroxy acids such as lactic acid and glycolic acid and their 15 derivatives and salts; or beta-hydroxy acids such as salicylic acid and salicylic acid salts and derivatives), urea, hyaluronic acid, phytic acid, lipoic acid; lysophosphatidic acid, skin peel agents (e.g., phenol, resorcinol and the like), vitamin B3 compounds (e.g., niacinamide, nicotinic acid and nicotinic acid salts and esters, including non-vasodilating esters of nicotinic acid (such as tocopheryl 20 nicotinate), nicotinyl amino acids, nicotinyl alcohol esters of carboxylic acids, nicotinic acid N-oxide and niacinamide N-oxide), vitamin B5 and retinoids (e.g., retinol, retinal, retinoic acid, retinyl acetate, retinyl palmitate, retinyl ascorbate). In the case of dry, scaly skin (xerosis) and ichthyosis such agents can alleviate the symptoms by temporary relief of itching associated with these conditions.

25 **Anti-oxidants/radical scavengers**

A safe and effective amount of an anti-oxidant/radical scavenger may be added to the compositions of the subject invention, preferably from about 0.1% to about 10% (w/w), more preferably from about 1% to about 5% (w/w), of the composition.

30 Anti-oxidants/radical scavengers such as ascorbic acid (vitamin C) and its salts, ascorbyl esters of fatty acids, ascorbic acid derivatives (e.g., magnesium ascorbyl phosphate, sodium ascorbyl phosphate, ascorbyl sorbate), tocopherol

(vitamin E), tocopherol sorbate, tocopherol acetate, other esters of tocopherol, butylated hydroxy benzoic acids and their salts, 6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid (commercially available under the tradename Trolox.sup.R), gallic acid and its alkyl esters, especially propyl gallate, 5 uric acid and its salts and alkyl esters, sorbic acid and its salts, lipoic acid, amines (e.g., N,N-diethylhydroxylamine, amino-guanidine), sulphhydryl compounds (e.g., glutathione), dihydroxy fumaric acid and its salts, lycine pidolate, arginine pilolate, nordihydroguaiaretic acid, bioflavonoids, curcumin, lysine, methionine, proline, superoxide dismutase, silymarin, tea extracts, grape skin/seed extracts, 10 melanin, and rosemary extracts may be used.

The foam of the present invention is suitable for delivering skin protecting and revitalizing anti-oxidants/radical scavengers. It is further pointed out that polyunsaturated fatty acids, containing omega-3 and omega-6 fatty acids (e.g., linoleic and linolenic acid, gamma-linoleic acid (GLA), eicosapentaenoic acid 15 (EPA) and docosahexaenoic acid (DHA) are beneficial in the treatment of psoriasis and other skin inflammation conditions. Likewise, emollients and silicone oils exert moisture-retaining and skin protective effects on the skin. Thus in a preferred embodiment, a skin protective foam is provided, wherein the hydrophobic solvent comprises in full or in part, a solvent, selected from the 20 group of emollients, silicone oil and oils, rich in unsaturated fatty acids, thus, affording a synergistic therapeutic effect of the anti-oxidants/radical scavenger agent and the vehicle components.

Self-tanning active agents

The foam of the present invention is particularly suitable for the uniform 25 delivery of a tanning active agent onto large areas of the skin. It is preferable that the compositions contain from about 0.1% to about 20%, more preferably from about 2% to about 7%, and still more preferably from about 3% to about 6%, of the composition, of dihydroxyacetone, or any other compound, known in the art as an artificial tanning active agent.

30 Skin Lightening and Whitening Agents

The foam of the present invention is particularly suitable for the uniform delivery of a skin lightening agent. When used, the compositions preferably

contain from about 0.1% to about 10%, more preferably from about 0.2% to about 5%, of the composition, of a skin-lightening agent. Suitable skin lightening or whitening agents include those known in the art, including hydroquinone, azelaic acid and other related dicarboxylic acids, and salts and derivatives thereof,

5 retinoids, kojic acid, arbutin, nicotinic acid and its precursors, salts and derivatives, arbutin, ascorbic acid and salts and derivatives thereof (e.g., magnesium ascorbyl phosphate or sodium ascorbyl phosphate), and herbal extracts (e.g., licorice extract, mulberry extract, placental extract).

In one or more embodiments of the present invention, the foam
10 composition comprises a combination of a skin whitening agent and a sunscreen agent.

In one or more embodiments of the present invention, the foam
composition comprises a combination of a skin whitening agent and an inorganic
sunscreen agent. When inorganic sunscreen agents, e.g. TiO_2 , are rubbed onto
15 the skin, they leave a white coating, which provides an immediate (although
transient) whitening effect, which is highly desirable by the consumer, who
wishes to see instant change in his/her appearance. The whitening agent, in
combination with the inorganic sunscreen agent in the foam carrier can be easily
and uniformly distributed on the skin surface, thereby affording an even instant
20 whitening effect, unlike creams that are difficult to spread evenly on skin areas.

Sunscreens

Exposure to ultraviolet light can result in excessive scaling and texture
changes of the stratum corneum. The foam of the present invention is
advantageous for the delivery of sunscreen agents. Its application is very
25 convenient and it spreads easily over large skin areas. The presence of a
hydrophobic solvent in the foam ensures long lasting effect, even while bathing.

As used herein, "sunscreen active" or "sunscreen agent" includes both
sunscreen agents and physical sunblocks. Suitable *sunscreen* actives may be
organic or inorganic.

30 Inorganic sunscreens useful herein include the following metallic oxides;
titanium dioxide having an average primary particle size of from about 15 nm to
about 100 nm, zinc oxide having an average primary particle size of from about

15 nm to about 150 nm, zirconium oxide having an average primary particle size of from about 15 nm to about 150 nm, iron oxide having an average primary particle size of from about 15 nm to about 500 nm, and mixtures thereof. When used herein, the inorganic sunscreens are present in the amount of from about 5 0.1% to about 20%, preferably from about 0.5% to about 10%, more preferably from about 1% to about 5%, of the composition.

A wide variety of conventional organic sunscreen actives are suitable for use herein. Specific suitable sunscreen actives include, for example: p-aminobenzoic acid, its salts and its derivatives (ethyl, isobutyl, glyceryl esters; p-dimethylaminobenzoic acid); anthranilates (i.e., o-amino-benzoates; methyl, menthyl, phenyl, benzyl, phenylethyl, linalyl, terpinyl, and cyclohexenyl esters); salicylates (amyl, phenyl, octyl, benzyl, menthyl, glyceryl, and di-pro-pyleneglycol esters); cinnamic acid derivatives (menthyl and benzyl esters, a-phenyl cinnamonnitrile; butyl cinnamoyl pyruvate); dihydroxycinnamic acid derivatives (umbelliferone, methylumbelliferone, methylaceto-umbelliferone); trihydroxy-cinnamic acid derivatives (esculetin, methylesculetin, daphnetin, and the glucosides, esculin and daphnin); hydrocarbons (diphenylbutadiene, stilbene); dibenzalacetone and benzalacetophenone; naphtholsulfonates (sodium salts of 2-naphthol-3,6-disulfonic and of 2-naphthol-6,8-disulfonic acids); di-20 hydroxynaphthoic acid and its salts; o- and p-hydroxybiphenyldisulfonates; coumarin derivatives (7-hydroxy, 7-methyl, 3-phenyl); diazoles (2-acetyl-3-bromoindazole, phenyl benzoxazole, methyl naphthoxazole, various aryl benzothiazoles); quinine salts (bisulfate, sulfate, chloride, oleate, and tannate); quinoline derivatives (8-hydroxyquinoline salts, 2-phenylquinoline); hydroxy- or 25 methoxy-substituted benzophenones; uric and violuric acids; tannic acid and its derivatives (e.g., hexaethylether); (butyl carbotol) (6-propyl piperonyl) ether; hydroquinone; benzophenones (oxybenzene, sulisobenzene, dioxybenzene, benzoresorcinol, 2,2',4,4'-tetrahydroxybenzophenone, 2,2'-dihydroxy-4,4'-dimethoxybenzophenone, octabenzene; 4-isopropylbenzoylmethane; 30 butylmethoxybenzoylmethane; etocrylene; octocrylene; [3-(4'-methylbenzylidene bornan-2-one), terephthalylidene dicamphor sulfonic acid and 4-isopropyl-di-benzoylmethane.

A safe and effective amount of the organic sunscreen active is used, typically from about 1% to about 20%, more typically from about 2% to about 10% of the composition. Exact amounts will vary depending upon the sunscreen or sunscreens chosen and the desired Sun Protection Factor (SPF).

5 **Agents for Hair Growth Disorders**

Agents, which affect the pattern of hair growth, can be suitably incorporated in the foam of the present invention. Male pattern baldness (MPB), the commonest cause of balding, is induced by the activity of the male hormone dihydrotestosterone (DHT), which converted from the hormone testosterone by 10 the enzymes 5 alpha reductase. Current treatments of MPB include minoxidil and agents, which inhibit 5 alpha reductase, such as finasteride, spironolactone, azelaic acid and azelaic acid derivatives and salts. Such agents, as well as other agents known in the art, can be incorporated in the foam of the present invention.

It is further pointed out that polyunsaturated fatty acids, i.e., such which 15 include any of the essential fatty acids (EFA's): linoleic and linolenic acid, gamma-linoleic acid (GLA), eicosapentaenoic acid (EPA) and docosahexaenoic acid (DHA), are also known to contribute to hair growth. Thus in a preferred embodiment, a hair growth foam is provided, wherein the hydrophobic solvent comprises in full or in part, an oil, rich in such unsaturated fatty acids.

20 **Figure-forming Agents; Agents to Treat Cellulite / Slimming**

Figure forming agents such as used in the treatment of cellulite and in slimming products, can be suitably incorporated in the foam of the present invention. A non-limiting exemplary list of active agents, known in the treatment of cellulite and in the induction of a slimming effect include herbal extracts, such 25 as bladderwrack extract, butcher's broom, cayenne, dandelion, red clover, ginkgo biloba, horse chestnut, witch hazel and borage oil, omega 3 and omega 6 oils, caffeic acid and salts and derivatives thereof, xanthine agents, such as caffeine, theophylline and pentoxyphilline, and nicotinic acid and salts and derivatives thereof.

30 **Agents to Treat Sunburn, Heat Burn, Radiation Burn, Rash and Itch**

Cosmetic and pharmaceutical ingredients which are known in the art of pharmacology and cosmetology to treat dermatitis, minor skin irritations, sunburn,

heat burn, radiation burn, and inhibit inflammation can be beneficially incorporated in the foam of the present invention.

Examples of such active agents include chamomile extract (*matricaria recutititia*), cucumber distillate (*cucumis sativus*), lavender water (*lavendula angustifolia*), rose water (*rosa damascena*), witch hazel (*hamamelis virginiana*), 5 allantoin, bisabolol, rosehip oil, calendula oil, azulaene, menthol and camphor.

Use of the Foam as a Lubricating and Protective Foam

There are several potential uses of the foam, particularly the silicone-oil based foam, as a lubricating foam. Typical examples are shaving foam, moisture 10 protection foam and antifriction foam. For such purposes, the foam can be used in its basic composition (without additional formulation aids and active ingredients), or with the addition of such additives.

FOAM FOR NEUTRALIZATION AND/OR DECONTAMINATION OF HAZARDOUS CHEMICALS AND TREATMENT OF HEAT BURNS

15 It has been reported that povidone iodine antiseptic, a popular iodine product, can ameliorate damage to guinea pig skin exposed to mustard gas and other chemical irritants and further reduces, and many times prevents, damage to human skin after accidental heat burns caused by hot water, oil or hot steam.

20 Other active compound, having decontamination abilities, comprise strong oxidants and free radical liberating compounds, such as hydrogen oxide, bleaching agents (e.g., sodium, calcium or magnesium hypochloride and the like) iodine, chlorohexidine and benzoyl peroxide.

25 The alcohol-free foam of the present invention, comprising one or more of the above decontaminating and neutralizing agents can be applied onto the contaminated skin to form a preventive layer, prior to contamination measure or as a decontamination/neutralization means, right after contamination has occurred.

PENETRATION ENHancers

A penetration enhancer or permeation enhancer is an agent used to 30 increase the permeability of the skin to a pharmacologically active agent to increase the rate at which the drug diffuses through the skin and enters the tissues and bloodstream. A chemical skin penetration enhancer increases skin

permeability by reversibly altering the physiochemical nature of the stratum corneum to reduce its diffusional resistance. In a review of the technical and patent literature up to 1996, numerous chemical compounds were cited as skin penetration enhancers. Most of the compounds are generally recognized as safe

5 (GRAS) ingredients that would often be considered inert by a formulator (Osborne D W, Henke J J, *Pharmaceutical Technology*, November 1997, pp 58-86.)

Examples of penetration enhancers, according to the present invention include: polyols, such as propylene glycol, hexylene glycol, diethylene glycol, propylene glycol n-alkanols, terpenes, di-terpenes, tri-terpenes, terpen-ols, 10 limonene, terpene-ol, 1-menthol, dioxolane, ethylene glycol, other glycols, and glycerol; sulfoxides, such as dimethylsulfoxide (DMSO), dimethylformamide, methyl dodecyl sulfoxide, dimethylacetamide; monooleate of ethoxylated glycerides (with 8 to 10 ethylene oxide units); Azone (1-dodecylazacycloheptan-15 2-one), 2-(n-nonyl)-1,3-dioxolane; esters, such as isopropyl myristate/palmitate, ethyl acetate, butyl acetate, methyl propionate, capric/caprylic triglycerides, octylmyristate, dodecyl-myristate; myristyl alcohol, lauryl alcohol, lauric acid, lauryl lactate ketones; amides, such as acetamide oleates such as triolein; various surfactants, such as sodium lauryl sulfate; various alkanoic acids such as 20 caprylic acid; lactam compounds, such as azone; alkanols, such as oleyl alcohol; dialkylamino acetates, and admixtures thereof.

Lower alcohols, such as ethanol, propanol, isopropanol, butanol, iso-butanol, t-butanol and pentanol are not considered appropriate penetration enhancers according to the present invention, due to their skin drying and 25 irritation properties.

Yet, another preferred class of penetration enhancers in the cyclodextrines and related compounds. Cyclodextrins are structurally related cyclic oligomaltoses which form a new group of pharmaceutical excipients. These are torus-shaped molecules with a hydrophilic outer surface and a lipophilic central 30 cavity. Cyclodextrins are capable of forming water-soluble inclusion complexes with a wide variety of lipophilic water-insoluble drugs by taking up a whole drug molecule, or some part of it, into the cavity. The cyclodextrin molecules are

relatively large (molecular weight ranging from almost 1000 to over 1500), with a hydrated outer surface, and under normal conditions, cyclodextrin molecules will only permeate the skin barrier with considerable difficulty. It is generally believed that the cyclodextrin molecules act as true carriers by keeping lipophilic drug

- 5 molecules in solution and deliver them to the skin surface where they partition from the cyclodextrin cavity into the skin.

FURTHER TECHNICAL PARAMETERS

The composition of the present invention may be contained in and

- 10 dispensed from a container capable of withstanding the pressure of the propellant gas and having an appropriate valve/nozzle for dispensing the composition as foam under pressure. A customary liquefied or compressed gas propellant can be added, in the amount of about 5-25% of the total composition. Liquefied propellants are gases that exist as liquids under pressure, including high purity
- 15 hydrocarbons such as propane, isobutane and n-butane, dimethyl ether and chlorofluorocarbons (CFCs). Compressed gasses are exemplified by air, nitrogen and carbon dioxide.

A specific embodiment according to the present invention comprises placing the composition of the present invention on a patch, occlusive tape or the

- 20 skin-contact compartment of a transdermal delivery apparatus and applying such object onto the skin, in order to attain effective superficial treatment or enhanced penetration of the drug into the skin or through the skin.

Utilizing such strategy, one can apply drugs, which are currently administered systemically or that require transdermal delivery, in the preferred

- 25 therapeutic system of the present invention. Examples for such drugs are nicotine, testosterone and other male hormones and male hormone precursors, estrogen and other female hormones and hormone precursors, growth hormone, insulin, caffeine, steroid and non-steroidal antiinflammatory agents and thyroid hormone substitutes.
- 30 The general process, as typically exemplified in Example 1 may be applied in order to produce the composition of the present invention.

The pharmaceutical carrier according to the present invention can also be used to prepare cosmetics for beauty purpose by adding into skin care agents and perfume.

EXAMPLES

The invention is described with reference to the following examples. This invention is not limited to these examples and experiments. Many variations will 5 suggest themselves and are within the full intended scope of the appended claims.

Example 1 - General Procedure for Preparing Foamable Composition.

Aqueous Phase: Water gelling agent and surface-active agent are dissolved in water, with agitation. The solution is warmed to 50-70⁰C. Water 10 soluble cosmetic or pharmaceutical active ingredients and optional water soluble ingredients are added with agitation to the Aqueous Phase mixture.

Hydrophobic Phase: The hydrophobic solvent is heated to same temperature. Foam adjuvant agent is added to preheated hydrophobic solvent. Oil soluble cosmetic or pharmaceutical active ingredients* and optional oil soluble 15 formulation ingredients are added with agitation to the Hydrophobic Phase mixture.

The warm Hydrophobic Phase is gradually poured into the warm Aqueous Phase, with agitation, followed by Ultraturax homogenization. The mixture is allowed to cool down to ambient temperature. In case of heat sensitive active 20 ingredients, the active ingredient is added with agitation to the mixture after cooling to ambient temperature. The mixture, at ambient temperature, is added to an aerosol container, the container is sealed and appropriate amount of propellant (5-25 w% of the composition mass) is added under pressure into the container.

Example 2 – Vegetable Oil-Based Foam Carrier Composition

Ingredient	Version	Version	Version	
	No. 1	No. 2	No. 3	
% (W/W)				
Hydrophobic solvent	Soybean oil	40	30.5	20
Water	Water	48.5	32.5	61
Foam adjuvant agent	Stearyl Alcohol	0.8	1.05	0.73
Surface-active agent	Sucrose ester SP70	0.64	0.45	0.8
Water gelling agent	Xanthan Gum	0.16	0.11	0.1
	Methocel ELV15	0.32	0.22	0.28
Other Ingredients	Antioxidant	0.02	0.02	0.02
	Preservatives	0.3	0.3	0.3
	Fragrance	0.2	0.2	0.2
Foam Specific gravity (gr/mL)		0.10	0.15	0.065

The compositions use a non-ionic surfactant and contain a combined amount of surface-active agent, foam adjuvant and water gelling agent ranging from 1.83 % to 1.92 % (w/w). The foam of this example is useful as a carrier of active pharmaceutical and/or cosmetic active ingredients, as exemplified below. It also can be used as a protective product. Additionally, it is also useful as lubricating foam, for various purposes.

Example 3 – Silicone Oil-Based Foam Carrier Composition

Specific Ingredient	Version	Version	
	No. 1	No. 2	
% (W/W)			
Hydrophobic solvent	Dimeticone 350*	25	10
Water	Water	72	87
Foam adjuvant agent	Stearyl Alcohol	0.2	0.2
Surface-active agent	Sucrose ester SP70	0.8	-
	Myrj 49P	-	0.8
Water gelling agent	Xanthan Gum	0.2	0.2
	Methocel ELV15	0.4	0.4
Other Ingredients	Antioxidant	0.02	0.02
	Preservatives	1	1
	Fragrance	0.2	0.2
Foam Specific gravity (gr/mL)		0.10	ND

* Dimethylpolysiloxane of 350 cps viscosity.

5 The compositions use only non-ionic surfactant and contain a combined amount of surface-active agent, foam adjuvant and water gelling agent of 1.6% (w/w). The foam of this example is useful as a carrier of active pharmaceutical and/or cosmetic active ingredients, as exemplified below. It also can be used as a protective product. Additionally, it is also useful as lubricating foam, for various

10 purposes.

Example 6 – Mineral Oil-Based Foam Carrier Composition

Ingredient	Version	Version	Version	Version	Version
	No. 1	No. 2	No. 3	No. 4	No. 5
% (W/W)					
Hydrophobic solvent	Mineral oil	69	50	50	25
Water	Water	28.4	46.7	46.7	71.88
Foam adjuvant agent	Stearyl Alcohol	0.7	1	1	0.5
Surface-active agent	Sucrose ester SP70	0.4	0.64	0	0.8
	PEG S-40	0	0	0.64	0
	Polysorbate-60	0	0	0	0.8
Water gelling agent	Xanthan Gum	0.1	0	0.14	0.2
	Methocel ELV15	0.2	0.4	0.32	0.4
Other Ingredients	Antioxidant	0.02	0.02	0.02	0.02
	Preservatives	1	1	1	1
	Fragrance	0.2	0.2	0.2	0.2
Foam Specific gravity (gr/mL)		ND	ND	ND	0.1

The compositions use only non-ionic solvents, and the total amount of surface active agent, foam adjuvants and water gelling agents ranges from 1.4 to 2.1 % (w/w). The foam of this example is useful as a carrier of active pharmaceutical and/or cosmetic active ingredients, as exemplified in examples below. It is also useful as lubricating foam, for various purposes.

Example 7 – Mixed Oils Foam Carrier Composition

	Ingredient	Version No. 1 25% Oil	Version No. 2 12.5% Oil
Hydrophobic solvent	Mineral oil	11.2%	5.6%
	Isopropyl myristate	5.0%	2.5%
	MCT oil	7.5%	3.8%
Foam adjuvant agent	Stearyl Alcohol	0.5%	0.25%
Water	Water	73.0%	85.2%
Surface-active agent	Sucrose ester SP70	0.8%	0.8%
	Distilled monoglyceride	1.2%	0.6%
Water gelling agent	Sodium lauryl sulphate	0.1%	0.1%
	Xanthan Gum	0.3%	0.3%
	Methocel ELV15	0.6%	0.6%

The foams of this example have a non-ionic surfactant to ionic surfactant ratio (w/w) of 20:1 and 14:1 for versions 1 and 2, respectively. Total amounts of surface active agent foam adjuvant and water gelling agent is in the range of 1.75 – 3.5 % (w/w). It is useful as a carrier of active pharmaceutical and/or cosmetic active ingredients, as exemplified in examples below. It is also useful as lubricating foam, for various purposes.

The following examples, representing optional drug-containing foams, are prototype formulations, which have not been optimized for stability and inter-component compatibility. Such optimization is a customary need, which can be done, using means, known to those skilled in the art of pharmaceutical

5 formulation

Example 8 – Antibacterial Foam Composition

Ingredient	Version 1 "Mupirocin"	Version 2 "Triple Antibiotic"	Version 3 "Fucidic Acid"	Version 4 "Metro- nidazole"	Version 5 "Triple Antibiotic"
Carrier Ingredients					
Mineral oil	48.8%	11.2%	48.8%	5.6%	5.6%
Isopropyl myristate		5.0%		2.5%	2.5%
MCT oil		7.5%		3.8%	3.8%
Stearyl Alcohol	0.8%	0.5%	0.8%	0.25%	0.25%
Water	50%	73.0%	50%	85.2%	85.2%
Sucrose ester SP70	0.8%	0.8%	-		0.8%
Myrj 40	-		0.8%	-	-
Distilled monoglyceride		1.2%		0.6%	0.6%
Tween 60				0.8%	
Sodium lauryl sulphate	0.05%	0.1%	0.1%		
Xanthan Gum	0.2%	0.3%	0.2%	0.3%	0.3%
Methocel ELV15	0.2%	0.6%	0.2%	0.6%	0.6%
Active Ingredients					
Mupirocin	2%				
Polymyxin B Sulfate		10,000 Units/gr			10,000 Units/gr
Bacitracin Zinc		500 Units/gr			500 Units/gr
Neomycin Sulfate*		0.05%			0.05%
Pramoxine HCl		1%			1%
Fucidic acid			2%		
Metronidazole				1%	

10 The foams of this example contain 100% non-ionic surfactant or have a non-ionic surfactant to ionic surfactant ratio ranging from 20:1 to 8:1. Total amounts of surface active agent, foam adjuvant and water gelling agent ranges from 2.05 – 3.5 % (w/w). It is useful for the treatment of bacterial skin infection

(general), cellulites, open wounds, cutaneous abscesses, furuncles, insect bite, impetigo, acne, acne-rosacea, and trichomonas vaginitis.

In certain embodiments, the foam of this example is useful for the prevention, decontamination and/or neutralization hazardous bacterial infestation
 5 (such as warfare organisms).

Example 9 – Antifungal Foam Composition

Ingredient	Version 1 "Terbinafine"	Version 2 "Clotrimazole"	Version 3 "Nystatin"	Version 4 "Nystatin"
Carrier Ingredients				
Mineral oil	48.8%	11.2%	48.8%	5.6%
Isopropyl myristate		5.0%		2.5%
MCT oil		7.5%		3.8%
Stearyl Alcohol	0.8%	0.5%	0.8%	0.25%
Water	50%	73.0%	50%	85.2%
Sucrose ester SP70	0.8%	0.8%	-	0.8%
Myrj 40	-		0.8%	-
Tween 80				0.8%
Distilled monoglyceride		1.2%		0.6%
Sodium lauryl sulphate	0.05%	0.1%	0.1%	
Xanthan Gum	0.2%	0.3%	0.2%	0.3%
Methocel ELV15	0.2%	0.6%	0.2%	0.6%
Active Ingredients				
Terbinafine	1%			
clotrimazole		2%		
Nystatin			100,000 Units/gr	100,000 Units/gr

10 The foams of this example have 100% non-ionic surfactant or have a non-ionic surfactant to ionic surfactant ratio ranging from 20:1 to 8:1. Total surface active agent, foaming adjuvant and water gelling agent ranges from 2.05 to 3.5% (w/w). It is useful in the treatment of dermatophyte infections, Tinea corporis, Tinea pedis, Tinea rubrum, Tinea unguium, Tinea cruris, Tinea barbae, and yeast
 15 infections, such as Candidiasis, Tinea versicolor and Candidal vaginitis.

Example 10 – Corticosteroid Foam Composition

Ingredient	Version 1 "Hydrocortisone"	Version 2 "Betamethasone"	Version 3 "Dexamethasone"
Carrier Ingredients			
Mineral oil	48.8%	11.2%	5.6%
Isopropyl myristate		5.0%	2.5%
MCT oil		7.5%	3.8%
Stearyl Alcohol	0.8%	0.5%	0.25%
Water	50%	73.0%	85.2%
Sucrose ester SP70	0.8%	0.4%	0.8%
Tween 80		0.4%	
Distilled monoglyceride		1.2%	0.6%
Sodium lauryl sulphate	0.05%		0.1%
Xanthan Gum	0.2%	0.3%	0.3%
Methocel ELV15	0.2%	0.6%	0.6%
Active Ingredients			
Hydrocortisone	1%		
Betamethasone dipropionate		0.05%	
Dexamethasone acetate			0.1%

Ingredient	Version 4 "Triamcinolone"	Version 5 "Flumetasone"
Carrier Ingredients		
Mineral oil	48.8%	48.8%
Stearyl Alcohol	0.8%	0.8%
Water	50%	50%
Sucrose ester SP70	0.8%	0.8%
Sodium lauryl sulphate	0.05%	0.05%
Xanthan Gum	0.2%	0.2%
Methocel ELV15	0.2%	0.2%
Active Ingredients		
Triamcinolone acetonide	0.1%	
Flumetasone pivalate		0.02%

5 The foams of this example have either 100% non-ionic surfactant or have a non-ionic surfactant to ionic surfactant ratio ranging from 20:1 to 16:1. Total surface active agent, foaming adjuvant and water gelling agent ranges from 2.05 to 3.5% (w/w). Indications include psoriasis, contact dermatitis, atopic dermatitis,

seborrheic dermatitis, nummular dermatitis, inflammatory acne, chronic dermatitis of the hands and feet, generalized exfoliative dermatitis, stasis dermatitis, lichen simplex chronicus, herpes gestationis and pruritic urticarial papules and plaques of pregnancy.

5

Example 11 – Antiviral Foam Composition

Ingredient	Version 1 "Acyclovir"	Version 2 "Acyclovir"	Version 3 "α-Interferon"
Carrier Ingredients			
Mineral oil	48.8%	11.2%	5.6%
Isopropyl myristate		5.0%	2.5%
MCT oil		7.5%	3.8%
Stearyl Alcohol	0.8%	0.5%	0.25%
Water	50%	73.0%	85.2%
Sucrose ester SP70		0.8%	0.8%
Tween 80	0.8%		
Distilled monoglyceride		1.2%	0.6%
Sodium lauryl sulphate			0.1%
Xanthan Gum	0.2%	0.3%	0.3%
Methocel ELV15	0.2%	0.6%	0.6%
Active Ingredients			
Acyclovir	5%	5%	
α-Interferon			105 IU/g

The foams of this example have either 100% non-ionic surfactant or have

10 a non-ionic surfactant to ionic surfactant ratio of 14:1. Total surface active agent, foaming adjuvant and water gelling agent ranges from 2.05 to 3.5% (w/w). Indications include Herpes simplex, Herpes zoster, Herpes gestationis and Herpes simplex genital ulcers.

Example 12 – Insect repellent Foam Composition

Ingredient	%
Isopropyl myristate	2.0%
MCT oil	2.0%
Stearyl Alcohol	1.2%
Water	64.0%
Sucrose ester SP70	0.8%
Sodium lauryl sulphate	0.1%
Xanthan Gum	0.3%
Methocel ELV15	0.6%
Propylene glycol	15%
DEET	15%

5 **Example 14 – Comparative Tolerability and Acceptability Study of a Corticosteroid Foam Composition Vs. a Conventional Ointment**

A panel of eight testers was requested to apply about 0.5 gr. of the foam preparation of example 10, Version 2 on one arm and 0.5 gr. of commercial Betamethasone valerate ointment, in a double blind fashion. They were asked to 10 describe their feeling about the ease of application, ease of spreading, spreadability and penetrability of each of the products and to give their general rating for each of the products on a scale of 0-3 (0 = poor; 1=barely acceptable; 2=acceptable and 3=excellent).

As demonstrated in the following table, the foam preparation of example 15 10, Version 2 obtained higher rates in all aspects of the test.

Property	Foam Preparation	Commercial Betamethasone Valerate Ointment
	Mean Rating	Mean Rating
Ease of application	2.3	1.6
Ease of spreading	2.5	1.9
Spreadability	2.9	1.2
Penetrability	2.0	1.5
Lack of sticky feeling	2.4	1.0
Lack of greasy feeling	2.2	1.0
Lack of shiny look	1.9	1.4
Overall rating	2.5	1.4

Example 15: Human Safety and Efficacy Study of a Corticosteroid

Composition in psoriasis patients

Two patients with mild to moderate psoriasis were administered topically a

5 Betamethasone 0.12% foam (example 10, Version 2) twice daily for two weeks. Both patients improved significantly, as manifested by clearance of the psoriatic plaques flattening of the thickened lesions. Figure 1 provides an exemplary response to treatment in the elbows of one of these patients. While betamethasone is known for its effect in psoriasis, such a beneficial effect after 14

10 days treatment is exceptional. The accelerated effect was attributed to the improved convenience and therefore, improved compliance.

Example 16: Human Safety and Efficacy Study of a Corticosteroid

Composition in psoriasis patients

Four patients with moderate to severe, disseminated atopic dermatitis

15 were administered topically a Betamethasone 0.12% foam (example 10, Version 2) twice daily for two weeks. All patients improved significantly, as manifested by complete clearance of all treated lesions. Figure 2 provides exemplary responses to treatment in different body areas, after 10 days of treatment. While betamethasone is known for its effect in atopic dermatitis, such a beneficial effect

20 after 10 days treatment is exceptional. The patients claimed that the use of the foam of the present invention was significantly more convenient than the corresponding cream and ointment. Thus, the accelerated effect was attributed to the improved convenience and therefore, improved compliance.

Example 17 – Foam Compositions with Urea

Component	% w/w			
Mineral oil	6.00	6.00	6.00	6.00
Isopropylmeristat	6.00	6.00	6.00	6.00
Glyceryl monostearate	0.50	0.50	0.50	0.50
Stearyl alcohol	0.20	0.20	0.20	1.00
Urea	10.00	10.00	10.00	10.00
Xantan gum	0.30	0.30	0.30	0.30
Methocel K100M	0.30	0.30	0.30	0.30
Myrj 52				3.00
TWEEN 80				1.00
Myrj 49p				3.00
TWEEN 60	1.00	1.00	1.00	
Cocamidopropylbetaein	0.50	0.50		
Phenonip	0.30	0.30	0.30	0.30
Water	to 100.0	to 100.0	to 100.0	to 100.0
Butane/propane	8.00	8.00	18.00	18.00
FoamQuality	E	E	E	E
Density	n/a	0.023	n/a	0.024

Example 18 – Compositions with various penetration enhancers

Component	% w/w				
Mineral oil	6.00	6.00	6.00	6.00	6.00
Isopropyl myristate	6.00	6.00	6.00	6.00	6.00
Glyceryl monostearate	0.50	0.50	0.50	0.50	0.50
Stearyl alcohol	1.00	1.00	1.00	1.00	1.00
Xantan gum	0.30	0.30	0.30	0.30	0.30
Methocel K100M	0.30	0.30	0.30	0.30	0.30
TWEEN 60	1.00				
TWEEN 80		1.00	1.00	1.00	1.00
MYRJ 49p	3.00	3.00	3.00	3.00	3.00
Propylen glycol		5.00			
Glycofurool			1.00	10.00	
Urea					10.00
Cocoamidopropylbethaine	0.50	0.50	0.50	0.50	0.50
Lidocain base	4.00	4.00	4.00	4.00	4.00
Phenonip	0.30	0.30	0.30	0.30	0.30
Water	to 100				
Butane/propane	8	8	8	16	10
Foam Quality	E	E	E	E	E
Density	0.020	0.018	0.019	0.019	0.018

Component	% w/w	% w/w
Isopropyl myristate	30.00	30.00
Glyceryl monostearate	0.50	0.50
Stearic acid	0.45	0.45
Xantan gum	0.30	0.30
Methocel K100M	0.30	0.30
TWEEN 80	1.00	1.00
MYRJ 49p	3.00	3.00
Cocoamidopropylbethaine	0.50	0.50
Transcutol p	20.00	20.00
Hydrophilic drug	Effective concentration	
Hydrophobic drug		Effective concentration
Phenonip	0.30	0.30
Water	to 100.0	to 100.0
Butane/propane	8.00	8.00
FoamQuality	E	E
Density	0.020	0.020

CLAIMS

What is claimed is:

1. An alcohol-free foamable pharmaceutical or cosmetic carrier, comprising:
 - 5 (a) a foamable composition comprising:
 - about 2-75% by weight of composition of a liquid, non-volatile hydrophobic solvent;
 - about 80-98% by weight of composition of water;
 - about 0.1% to 5% by weight of composition of a foam adjuvant agent
 - 10 selected from the group consisting of fatty alcohols, fatty acids, hydroxyl-substituted fatty alcohols, hydroxyl-substituted fatty acids, and fatty acids and fatty alcohols including at least one double bond in its carbon atom chain;
 - about 0.1% to 5% by weight of composition of a surface-active agent; and
 - about 0.1% to 5% by weight of composition of a water gelling agent.
 - 15 and
 - (b) a liquefied or compressed gas propellant in a container,
 - which upon release provides a breakable foam suitable for topical or mucosal administration.
- 20 2. The foamable carrier of claim 1, wherein the hydrophobic solvent comprises about 5-10% by weight of composition.
 3. The foamable carrier of claim 1, wherein the hydrophobic solvent comprises about 10-20% by weight of composition.
- 25 4. The foamable carrier of claim 1, wherein the hydrophobic solvent comprises about 20-75% by weight of composition.
 5. The foamable carrier of claim 1, wherein the hydrophobic solvent comprises a mixture of mineral oil and an emollient in a ratio between 2:8 and 8:2 on a weight basis.

6. The foamable carrier of claim 1, wherein at least 2% of the foamable composition is a silicone oil.
7. The foamable carrier of claim 1, wherein the surface-active agent is 5 selected from the groups of non ionic surfactants, cationic surfactants, amphoteric and zwitterionic surfactants.
8. The foamable carrier of claim 1, wherein the surface-active agent is a mixture of a non ionic surfactant and an anionic surfactant in a 20:1 to 1:1 ratio.
10
9. The foamable carrier of claim 1, wherein the surface-active agent is a mixture of a non-ionic surfactant and an ionic surfactant in a 100:1 to 6:1 ratio.
10. The foamable carrier of claim 1, wherein the surface-active agent consists 15 essentially of one or more non-ionic surfactants.
11. The foamable carrier of claim 1, wherein the surface-active agent has HLB value of more than 9.
- 20 12. The foamable carrier of claims 7, 8, 9, or 10, wherein the non-ionic surfactant comprises a sucrose ester.
13. The foamable carrier of claim 1, wherein the hydrophobic solvent is selected from the group comprising vegetable oils, marine oils, mineral oils, 25 emollients, silicone oils, plant-derived therapeutic oils and mixture thereof.
14. The foamable carrier of claim 1, wherein the combined amount of foam adjuvant agent, surface-active agent and water gelling agent is less than about 8% (w/w).

30

15. The foamable carrier of claim 1, wherein the combined amount of foam adjuvant agent, surface-active agent and water gelling agent is less than about 5% (w/w) of the foamable composition.

5 16. A pharmaceutical or cosmetic composition, comprising:

(a) a foamable composition comprising:

about 2-75% by weight of composition of a liquid, non-volatile hydrophobic solvent;

about 80-98% by weight of composition of water;

10 about 0.1% to 5% by weight of composition of a foam adjuvant agent selected from the group consisting of fatty alcohols, fatty acids, hydroxyl-substituted fatty alcohols, hydroxyl-substituted fatty acids, and fatty acids and fatty alcohols including at least one double bond in its carbon atom chain;

about 0.1% to 5% by weight of composition surface-active agent; and

15 about 0.1% to 5% by weight of composition water gelling agent;

(b) a therapeutically effective amount of an active agent;

and

(c) a liquefied or compressed gas propellant in a container, which upon release provides a breakable foam suitable for topical or

20 muscasol administration.

17. The pharmaceutical or cosmetic composition of claim 16, wherein the hydrophobic solvent comprises about 5-10% by weight of composition.

25 18. The pharmaceutical or cosmetic composition of claim 16, wherein the hydrophobic solvent comprises about 10-20% by weight of composition.

19. The pharmaceutical or cosmetic composition of claim 16, wherein the hydrophobic solvent comprises about 20-75% by weight of composition.

30 20. The pharmaceutical or cosmetic composition of claim 16, wherein the active agent is a drug.

21. The pharmaceutical or cosmetic composition of claim 16, wherein the active agent is a cosmetically effective agent.
- 5 22. The pharmaceutical or cosmetic composition of claim 16, wherein the hydrophobic solvent is selected from the group comprising vegetable oils, marine oils, mineral oils, emollient, silicone oils, plant-derived therapeutic oils and mixture thereof at any proportion.
- 10 23. The pharmaceutical or cosmetic composition of claim 16, further comprising excipients selected from the group consisting of antioxidants, humectants, flavoring, colorant and odorant agents.
- 15 24. The pharmaceutical or cosmetic composition of claim 16, wherein the hydrophobic solvent comprises a mixture of mineral oil and an emollient in a ratio between 2:8 and 8:2 on a weight basis.
- 20 25. The pharmaceutical or cosmetic composition of claim 16, wherein at least 2% of the composition is a silicone oil.
26. The pharmaceutical or cosmetic composition of claim 16, wherein the surface-active agent is a mixture of a non ionic surfactant and an anionic surfactant in a 20:1 to 1:1 ratio.
- 25 27. The pharmaceutical or cosmetic composition of claim 16, wherein the surface-active agent consists essentially of one or more non-ionic surfactants.
28. The pharmaceutical or cosmetic composition of claim 16, wherein the surface-active agent is a mixture of a non-ionic surfactant and an ionic surfactant
- 30 30 in a 100:1 to 6:1 ratio.

29. The pharmaceutical or cosmetic composition of claim 16, wherein the combined amount of foam adjuvant agent, surface active agent and water gelling agent is less than about 8% (w/w).

5 30. The composition of claim 16, further comprising an effective concentration of a penetration enhancer.

31. The pharmaceutical or cosmetic composition of claim 16, wherein the combined amount of foam adjuvant agent, surface active agent and water gelling 10 agent is less than about 5% (w/w).

32. The pharmaceutical or cosmetic composition of claim 20 wherein the drug is selected for the treatment of a disease, the etiology of which is bacterial, fungal, viral, parasitic, inflammatory, autoimmune, allergic, hormonal, malignant 15 and combinations thereof.

33. The pharmaceutical or cosmetic composition of claim 20, wherein the drug is selected for the treatment of a superficial condition.

20 34. The composition of claim 20, wherein the drug is selected for the treatment of a disorder of the skin, mucosal membrane, vagina and rectum.

35. The composition of claim 20, wherein the drug is selected for the treatment of a disorder, selected from the group of dermatosis, dermatitis, bacterial 25 infections, fungal infections, parasitic infections, viral infections, disorders of hair follicles and sebaceous glands, acne, rosacea, scaling papular diseases, benign tumors, malignant tumors, reactions to sunlight, bullous diseases, pigmentation disorders, disorders of cornification, pressure sores, disorders of sweating, inflammatory reactions, xerosis, ichthyosis, allergy, burn, wound, cut, and non- 30 dermatological disorders, which respond to transdermal delivery of said drug.

36. The composition of claim 20 wherein the drug is selected for the treatment of wounds, burns, cuts and ulcers.
37. The composition of claim 20 wherein the drug is antibacterial.
5
38. The composition of claim 20 wherein the drug is antifungal.
39. The composition of claim 20 wherein the drug is antiviral.
- 10 40. The composition of claim 20 wherein the active agent is an insecticide.
41. The composition of claim 20 wherein the active agent is an insect repellent.
- 15 42. The composition of claim 20, wherein the drug is an anti-inflammatory or antiallergic agent.
43. The composition of claim 20, wherein the drug is an anticancer agent.
- 20 44. The composition of claim 20, wherein the drug is a photodynamic therapy agent.
45. The composition of claim 20 wherein the drug is a local anesthetic.
- 25 46. The composition of claim 20 wherein the drug is a nonsteroidal anti-inflammatory drug (NSAID)
47. The composition of claim 21, wherein the active agent is a retinoid.
- 30 48. The composition of claim 21, wherein the active agent is an anti-wrinkle agent.

49. The composition of claim 20 or 21, wherein the active agent is a skin-whitening agent.

50. The composition of claim 20 or 21, wherein said active agent is selected
5 from the group comprising sulfur-containing amino acids, thiol compounds, alpha hydroxy acids, lactic acid and its derivatives and salts, glycolic acid and its derivatives and salts, beta-hydroxy acids, salicylic acid and salicylic acid salts and derivatives, phytic acid, lipoic acid, lysophosphatidic acid, skin peel agents, phenol, resorcinol, vitamin B3 compounds, niacinamide, nicotinic acid and
10 nicotinic acid salts and esters, tocopheryl nicotinate, nicotinyl amino acids, nicotinyl alcohol esters of carboxylic acids, nicotinic acid N-oxide and niacinamide N-oxide, retinoids, retinol, retinal, retinoic acid, retinyl acetate, retinyl palmitate, retinyl ascorbate, caffeine, theophiline, pentoxyphilline, dihydroxy acetone kojic acid, arbutin, nicotinic acid and its precursors, salts and derivatives, arbutin,
15 ascorbic acid and salts and derivatives thereof.

51. The composition of claim 20 or 21, wherein the active agent is an herbal extract.

20 52. The composition of claim 20 or 21, wherein the active agent is a radical scavenger.

53. The composition of claim 21, wherein the active agent is a self-tanning agent.

25 54. The composition of claim 20 or 21, wherein the active agent is an anti-acne active agent.

55. The composition of claim 20 or 21, wherein the active agent is a skin
30 whitening agent.

56. The composition of claim 20 or 21 wherein the active agent is a figure forming agent.

57. The composition of claim 21, wherein the active agent is an agent that influences hair growth.

58. The composition of claim 21, wherein the active agent is a hair growth stimulating agent.

10 59. The composition of claim 21, wherein the active agent is a hair growth inhibiting agent.

60. The composition of claim 16, further comprising a sunscreen agent.

15 61. The composition of claim 16, further comprising an inorganic sunscreen agent.

62. The composition of claim 16, wherein the active agent is a combination of a skin whitening agent and a sunscreen agent.

20 63. The composition of claim 16, wherein the active agent is a combination of a skin whitening agent and an inorganic sunscreen agent.

64. The composition of claim 20 wherein the drug is intended for transdermal delivery.

25 65. The composition of claim 16, further comprising a decontaminating agent selected from the group comprising an oxidizing agent, iodine and iodine compounds, chlorohexidine, bleaching agents and surface-active agents.

30 66. A method of treating, alleviating or preventing a dermatological disorder, comprising:

administering topically to a subject having said dermatological disorder a therapeutically effective amount of a breakable foam composition comprising:

- (a) a foamable composition comprising:
 - 5 about 2-75% by weight of composition of a liquid, non-volatile hydrophobic solvent;
 - about 80-98% by weight of composition of water;
 - about 0.1% to 5% by weight of composition of a foam adjuvant agent selected from the group consisting of fatty alcohols, fatty acids, hydroxyl-
 - 10 substituted fatty alcohols, hydroxyl-substituted fatty acids, and fatty acids and fatty alcohols including at least one double bond in its carbon atom chain;
 - about 0.1% to 5% by weight of composition of a surface-active agent; and
 - about 0.1% to 5% by weight of composition of a water gelling agent.
- 15 (b) at least one active agent, which is intended to prevent, alleviate or cure said disorder; and
- (c) a liquefied or compressed gas propellant.

20 67. The method of claim 66, wherein at least 2% of the composition is a silicone oil.

68. The method of claim 66, wherein the hydrophobic solvent comprises a mixture of mineral oil and an emollient in a ratio between 2:8 and 8:2 on a weight

25 basis.

69. The method of claim 66, wherein the surface-active agent is selected from the groups of non-ionic surfactants, cationic surfactants, amphoteric and zwitterionic surfactants.

30 70. The method of claim 66, wherein the surface-active agent is a mixture of a non-ionic surfactant and an anionic surfactant in a 1:1 to 20:1 ratio.

71. The method of claim 66, wherein the surface-active agent is non-ionic.
72. The method of claim 66, wherein the surface-active agent has HLB value
5 of more than 9.
73. The method of claim 66, wherein the non-ionic surfactant comprises a sucrose ester.
- 10 74. The method of claim 66, wherein the drug is intended for the treatment of a disease, the etiology of which is bacterial, fungal, viral, parasitic, inflammatory, autoimmune, allergic, hormonal, malignant and combinations thereof.
- 15 75. The method of claim 66, wherein the drug is selected from the group comprising antibacterial, antifungal, anti-inflammatory, antiallergic, nonsteroidal anti-inflammatory, retinoid, alpha hydroxy acid, beta hydroxy acid, keratolytic, antiproliferative, anticancer and anti-pigmentation drugs.
- 20 76. The method of claim 66, wherein the drug is selected from the group comprising an insecticide of an insect repellent.
77. The method of claim 66, wherein said compound is applied topically to an affected area.
- 25 78. The method of claim 66, wherein said dermatological disorder comprises dermatosis, dermatitis, bacterial Infections, fungal Infections, parasitic infections, viral infections, disorders of hair follicles and sebaceous glands, scaling papular diseases, benign tumors, malignant tumors, reactions to sunlight, bullous diseases, pigmentation disorders, disorders of cornification, pressure sores,
- 30 disorders of sweating, inflammatory reactions, xerosis, ichthyosis, allergy, burn, wound, cut, and non-dermatological disorders, which respond to transdermal delivery of said active agent.

79. The method of claim 66, wherein the active agent is a figure forming agent.

5 80. The method of claim 66, wherein the active agent is an agent that influences hair growth.

81. The method of claim 66, wherein the active agent is a hair growth stimulating agent.

10 82. The method of claim 66, wherein the active agent is a hair growth inhibiting agent.

83. The method of claim 66, further comprising a sunscreen agent.

15 84. The method of claim 66, further comprising an inorganic sunscreen agent.

85. The method of claim 66, wherein the active agent is a combination of a skin whitening agent and a sunscreen agent.

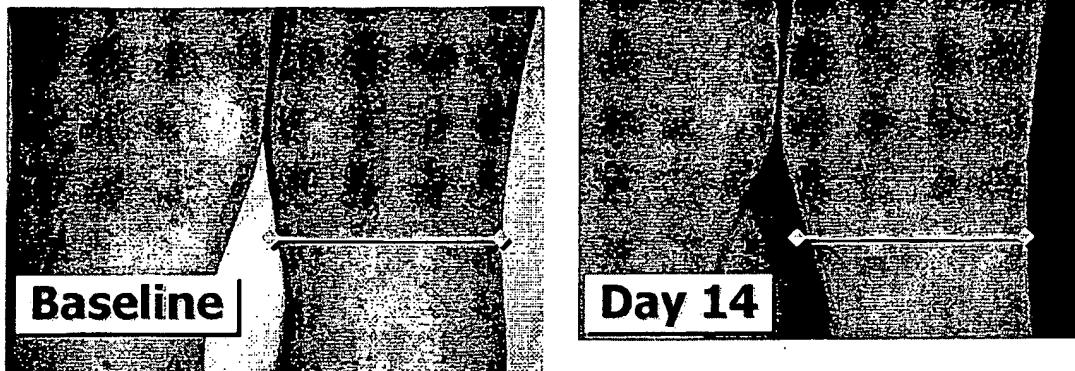
20 86. The method of claim 66, wherein the active agent is a combination of a skin whitening agent and an inorganic sunscreen agent.

87. The method of claim 66, comprising an effective concentration of a

25 penetration enhancer.

Picture No. 2: Improvement in the treatment of atopic dermatitis, using Bethasone 0.12% foam

**Patient No. 1, Atopic Dermatitis (Posterior part of the knee);
Betamethasone 0.12%**



Patient No. 2, Atopic Dermatitis (Arms); Betamethasone 0.12%

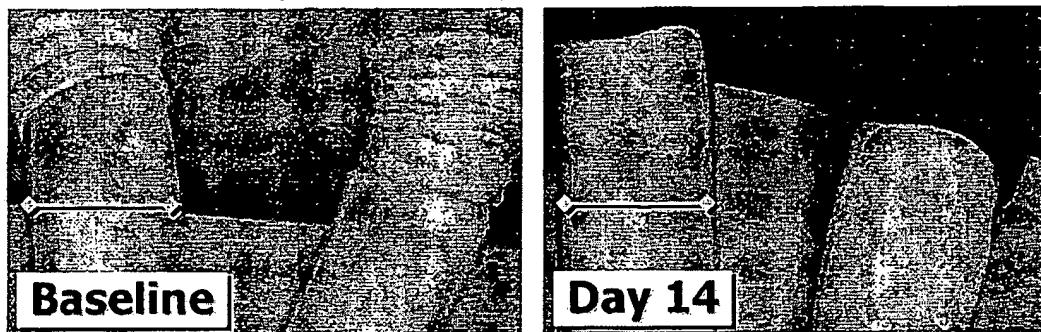


Figure 1

**Picture No. 1: Improvement in the treatment of psoriasis, using Bethasone
0.12% foam**

Patient No. 4, Psoriasis (Elbow)
Betamethasone 0.12%

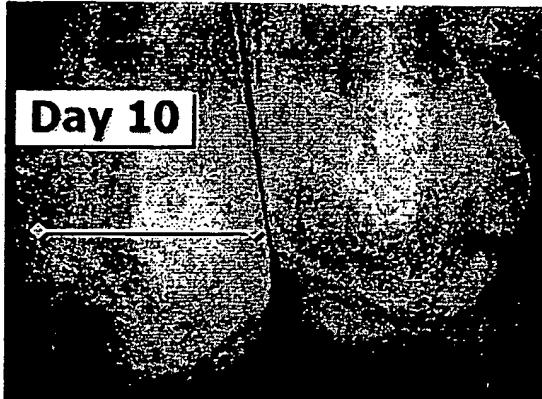
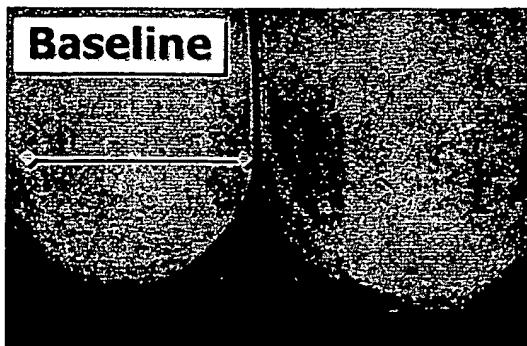


Figure 2

DUPLICATE

INTERNATIONAL SEARCH REPORT

25 APR 2005
101532618
International Application No
IB 03/05527

A. CLASSIFICATION OF SUBJECT MATTER
IPC 7 A61K9/12 A61K7/00 A01N25/16 A61K47/24

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
IPC 7 A61K A01N

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, EMBASE

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 03/051294 A (MEDICIS PHARMACEUTICALS) 26 June 2003 (2003-06-26) example 1 -----	1-87
X	US 5 326 557 A (MALCZWESKI REGINA M ET AL) 5 July 1994 (1994-07-05) See table II the whole document -----	1-87
X	US 5 435 996 A (MALCZWESKI REGINA M ET AL) 25 July 1995 (1995-07-25) the whole document -----	1-87
X	US 2002/072544 A1 (MILLER DENNIS ET AL) 13 June 2002 (2002-06-13) examples ----- -/-	1-87

 Further documents are listed in the continuation of box C. Patent family members are listed in annex.

* Special categories of cited documents :

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- *E* earlier document but published on or after the International filing date
- *L* document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- *O* document referring to an oral disclosure, use, exhibition or other means
- *P* document published prior to the international filing date but later than the priority date claimed

- *T* later document published after the International filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- *X* document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- *Y* document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- *&* document member of the same patent family

Date of the actual completion of the international search

5 July 2004

Date of mailing of the international search report

28.10.2004

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Giménez Miralles, J

INTERNATIONAL SEARCH REPORT

National Application No

PCT/IB 03/05527

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 6 093 408 A (HASENOEHRL ERIK JOHN ET AL) 25 July 2000 (2000-07-25) the whole document	1-87
X	----- US 5 922 331 A (MAUSNER JACK) 13 July 1999 (1999-07-13) the whole document	1-87
X	----- US 5 641 480 A (VERMEER ROBERT) 24 June 1997 (1997-06-24) the whole document	1-87
X	----- US 2002/098215 A1 (SIMONNET JEAN-THIERRY ET AL) 25 July 2002 (2002-07-25) the whole document	1-87
X	----- US 2002/035046 A1 (TAHAR MAURICE ET AL) 21 March 2002 (2002-03-21) the whole document	1-87
Y	----- WO 91/11991 A (KABI PHARMACIA AB) 22 August 1991 (1991-08-22) the whole document	1-87
Y	----- US 6 423 323 B2 (NEUBOURG FRITZ) 23 July 2002 (2002-07-23) cited in the application the whole document	1-87
Y	----- US 5 536 743 A (BORGMAN ROBERT J) 16 July 1996 (1996-07-16) cited in the application the whole document	1-87
Y	----- US 5 679 324 A (LISBOA LOUIS SERGIO ET AL) 21 October 1997 (1997-10-21) cited in the application the whole document	1-87
A	----- US 5 034 220 A (BURLANT WILLIAM J ET AL) 23 July 1991 (1991-07-23) column 7	1-87

INTERNATIONAL SEARCH REPORT

International Application No

/IB 03/05527

Patent document cited in search report		Publication date		Patent family member(s)		Publication date
WO 03051294	A	26-06-2003	US	2002164381 A1		07-11-2002
			CA	2470582 A1		26-06-2003
			WO	03051294 A2		26-06-2003
US 5326557	A	05-07-1994	US	5435996 A		25-07-1995
US 5435996	A	25-07-1995	US	5326557 A		05-07-1994
US 2002072544	A1	13-06-2002	DE	10035930 A1		31-01-2002
			EP	1174180 A1		23-01-2002
			JP	2002128635 A		09-05-2002
US 6093408	A	25-07-2000		NONE		
US 5922331	A	13-07-1999		NONE		
US 5641480	A	24-06-1997		NONE		
US 2002098215	A1	25-07-2002	FR	2804015 A1		27-07-2001
			BR	0100424 A		25-09-2001
			CA	2332469 A1		21-07-2001
			EP	1120102 A2		01-08-2001
			JP	2001226221 A		21-08-2001
US 2002035046	A1	21-03-2002	AU	9735901 A		27-06-2002
			CA	2365818 A1		21-06-2002
			CN	1366874 A		04-09-2002
			EP	1216685 A2		26-06-2002
			JP	2002322045 A		08-11-2002
			AU	5764800 A		22-01-2001
			WO	0101949 A1		11-01-2001
			US	2004167046 A1		26-08-2004
WO 9111991	A	22-08-1991	AT	151982 T		15-05-1997
			DE	69125825 D1		28-05-1997
			EP	0514465 A1		25-11-1992
			FI	923548 A		07-08-1992
			JP	5504571 T		15-07-1993
			NO	923078 A		28-09-1992
			WO	9111991 A1		22-08-1991
US 6423323	B2	03-01-2002	AT	210954 T		15-01-2002
			AU	9162498 A		08-03-1999
			CA	2300054 A1		25-02-1999
			DE	29824538 U1		16-08-2001
			DE	29824900 U1		15-05-2003
			DE	29825037 U1		29-04-2004
			DE	59802564 D1		31-01-2002
			DK	1014916 T3		15-04-2002
			WO	9908649 A2		25-02-1999
			EP	1014916 A2		05-07-2000
			ES	2170522 T3		01-08-2002
			JP	2001515022 T		18-09-2001
			PT	1014916 T		28-06-2002
			US	2003059394 A1		27-03-2003
			US	2002001599 A1		03-01-2002
US 5536743	A	16-07-1996	US	4837378 A		06-06-1989

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

Continuation of Box I.1

Rule 39.1(iv) PCT - Method for treatment of the human or animal body by therapy

Continuation of Box I.2

Claims Nos.: 1,16,66 partially

Present independent claim 1 relates to an extremely large number of possible foamable carrier compositions, namely all those comprising 1) a hydrophobic solvent, 2) water, 3) a foam adjuvant, 4) a surfactant, and 5) a gelling agent (with no restriction or limitation). Said definition covers so many possible options and alternatives lacking support and/or disclosure in the description (Articles 5 and 6 PCT) as to render a meaningful search impossible. It is the same for independent claims 16 and 66. Moreover, said foamable carrier composition is defined by reference to the result to be achieved, expressed in terms of a desirable characteristic or property, namely that upon release of a pharmaceutical or cosmetic composition comprising said foamable carrier composition from a pressurized container it produces a "breakable foam". The claims cover all foamable carrier compositions comprising the components 1)-5) and having this characteristic or property, whereas the application provides support within the meaning of Article 6 PCT and/or disclosure within the meaning of Article 5 PCT for only a very limited number of such compositions. In the present case, the claims so lack support, and the application so lacks disclosure, that a meaningful search over the whole of the claimed scope is impossible.

Further, the claims also lack clarity (Article 6 PCT), because the definition by reference to a result to be achieved only amounts to claiming the technical problem to be solved instead of claiming the relevant technical features which amount to its solution. Moreover, the expression "breakable foam" is unclear. Again, this lack of clarity in the present case is such as to render a meaningful search over the whole of the claimed scope impossible.

Consequently, the search has been carried out for those parts of the claims which appear to be clear, supported and disclosed, namely those parts relating to claims 1, 16 and 66 taken in combination with claims 6 and 12 and the examples.

The applicant's attention is drawn to the fact that claims relating to inventions in respect of which no international search report has been established need not be the subject of an international preliminary examination (Rule 66.1(e) PCT). The applicant is advised that the EPO policy when acting as an International Preliminary Examining Authority is normally not to carry out a preliminary examination on matter which has not been searched. This is the case irrespective of whether or not the

INTERNATIONAL SEARCH REPORT

International Application No

PCT/IB 03/05527

Patent document cited in search report	Publication date	Patent family member(s)	Publication date	
US 5536743	A	US 5840744 A	24-11-1998	
		AT 103176 T	15-04-1994	
		AU 635752 B2	01-04-1993	
		AU 5829490 A	07-01-1991	
		BR 9006793 A	06-08-1991	
		CA 1337279 C	10-10-1995	
		CN 1049101 A ,B	13-02-1991	
		DE 69007547 D1	28-04-1994	
		DE 69007547 T2	14-07-1994	
		DK 404376 T3	24-05-1994	
		EP 0404376 A2	27-12-1990	
		ES 2062366 T3	16-12-1994	
		HU 56490 A2	30-09-1991	
		IE 65413 B1	18-10-1995	
		IL 94619 A	24-01-1995	
		JP 2999820 B2	17-01-2000	
		JP 4500365 T	23-01-1992	
		KR 169487 B1	15-01-1999	
		PL 285507 A1	13-01-1992	
		RO 109156 B1	30-12-1994	
		RU 2032402 C1	10-04-1995	
		WO 9014832 A1	13-12-1990	
		ZA 9004413 A	24-04-1991	
		AT 103813 T	15-04-1994	
		AU 3043289 A	11-08-1989	
		AU 621589 B2	19-03-1992	
		CA 1297029 C	10-03-1992	
		CN 1035435 A ,B	13-09-1989	
		DE 68914365 D1	11-05-1994	
		DE 68914365 T2	11-08-1994	
		EP 0355152 A1	28-02-1990	
		JP 2503004 T	20-09-1990	
		JP 2714464 B2	16-02-1998	
		KR 135313 B1	23-04-1998	
		WO 8906537 A1	27-07-1989	
		ZA 8900233 A	26-09-1990	
US 5679324	A	21-10-1997	NONE	
US 5034220	A	23-07-1991	AU 7851191 A	07-01-1992
			CA 2082829 A1	21-12-1991
			EP 0581764 A1	09-02-1994
			WO 9119477 A1	26-12-1991

INTERNATIONAL SEARCH REPORT

International application No.
PCT/IB 03/05527

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
Although claims 66-87 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. Claims Nos.: **1,16,66 partially**
because they relate to parts of the International Application that do not comply with the prescribed requirements to such an extent that no meaningful International Search can be carried out, specifically:
see FURTHER INFORMATION sheet PCT/ISA/210
3. Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this International application, as follows:

see additional sheet

1. As all required additional search fees were timely paid by the applicant, this International Search Report covers all searchable claims.
2. As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. As only some of the required additional search fees were timely paid by the applicant, this International Search Report covers only those claims for which fees were paid, specifically claims Nos.:
4. No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:
1, 16, 66 partially

Remark on Protest

The additional search fees were accompanied by the applicant's protest.
 No protest accompanied the payment of additional search fees.

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

This International Searching Authority found multiple (groups of) inventions in this international application, as follows:

1. claims: 1,16,66 partially

A foamable carrier composition comprising 1) hydrophobic solvent, 2) water, 3) foam adjvant, 4) surfactant, and 5) gelling agent; and a pharmaceutical or cosmetic composition comprising said foamable carrier.

2. claims: 1,16,66 partially

A foam generator/dispenser (pressurized container) comprising a container and a liquified or compressed gas propellant

FURTHER INFORMATION CONTINUED FROM PCT/ISA/ 210

claims are amended following receipt of the search report or during any Chapter II procedure. If the application proceeds into the regional phase before the EPO, the applicant is reminded that a search may be carried out during examination before the EPO (see EPO Guideline C-VI, 8.5), should the problems which led to the Article 17(2) declaration be overcome.

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